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# PATENT COOPERATION TREATY

From the INTERNATIONAL BUREAU

PCT

## NOTIFICATION OF ELECTION

(PCT Rule 61.2)

To:

Commissioner  
US Department of Commerce  
United States Patent and Trademark  
Office, PCT  
2011 South Clark Place Room  
CP2/5C24  
Arlington, VA 22202  
ETATS-UNIS D'AMERIQUE

in its capacity as elected Office

Date of mailing (day/month/year)

11 June 2002 (11.06.02)

International application No.

PCT/IL01/00843

Applicant's or agent's file reference

11777/WO/00

International filing date (day/month/year)

06 September 2001 (06.09.01)

Priority date (day/month/year)

21 September 2000 (21.09.00)

Applicant

SONNENSCHNEIN, Elazar et al

1. The designated Office is hereby notified of its election made:



in the demand filed with the International Preliminary Examining Authority on:

01 April 2002 (01.04.02)



in a notice effecting later election filed with the International Bureau on:

2. The election ☒ was



was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

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## PATENT COOPERATION TREATY

## PCT

## INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference <b>11777/WO/00</b>	<b>FOR FURTHER ACTION</b> see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.	
International application No. <b>PCT/IL 01/ 00843</b>	International filing date (day/month/year) <b>06/09/2001</b>	(Earliest) Priority Date (day/month/year) <b>21/09/2000</b>
Applicant <b>MEDIGUS LTD.</b>		

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 5 sheets.

☒ It is also accompanied by a copy of each prior art document cited in this report.

## 1. Basis of the report

- a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

☐ the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

- b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :

☐ contained in the international application in written form.

☐ filed together with the international application in computer readable form.

☐ furnished subsequently to this Authority in written form.

☐ furnished subsequently to this Authority in computer readable form.

☐ the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

☐ the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. ☐ **Certain claims were found unsearchable** (See Box I).

3. ☒ **Unity of invention is lacking** (see Box II).

4. With regard to the **title**,

☒ the text is approved as submitted by the applicant.

☐ the text has been established by this Authority to read as follows:

5. With regard to the **abstract**,

☒ the text is approved as submitted by the applicant.

☐ the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No. \_\_\_\_\_

☐ as suggested by the applicant.

☐ because the applicant failed to suggest a figure.

☐ because this figure better characterizes the invention.

☒ None of the figures.

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# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/IL 01/00843

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
  
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
  
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☒ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
  
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
  
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
  
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☒ No protest accompanied the payment of additional search fees.

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**FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210**

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-6

Endoscope comprising two or more separate optical channels, where each optical channel comprises an objective lens for focusing the image on to an image sensor chip. Each objective lens is located at a different position along the length of the endoscope.

2. Claims: 7, 8-10

Endoscope and surgical stapler combination.

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## INTERNATIONAL SEARCH REPORT

International Application No

IL 01/00843

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 7 A61B1/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 91 00049 A (KARL STORZ GMBH & CO.) 10 January 1991 (1991-01-10) page 17, line 6 -page 23, line 14 figures 1-6C	1
A	---	2,4,5
X	DE 42 41 938 A (KARL STORZ GMBH & CO.) 16 June 1994 (1994-06-16) column 7, line 41 -column 9, line 33 figures	1
Y	---	7
Y	WO 00 53102 A (GAZAYERLI) 14 September 2000 (2000-09-14) page 7, line 20 -page 9, line 11 page 19, line 13 -page 23, line 3 figures 1-2B	7
A	---	8
	--- -/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

## \* Special categories of cited documents :

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

\*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

\*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*&\* document member of the same patent family

Date of the actual completion of the international search

21 June 2002

Date of mailing of the international search report

- 8. 07. 2002

Name and mailing address of the ISA

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Chen, A

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## INTERNATIONAL SEARCH REPORT

International Application No

IL 01/00843

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3 889 662 A (MITSUI) 17 June 1975 (1975-06-17) column 4, line 4 - line 50 ----	1,6
A	WO 89 11252 A (UNIVERSITY OF CONNECTICUT) 30 November 1989 (1989-11-30) page 19, line 3 -page 24, line 11 figures 1,2,4,5 ----	1-3
A	US 5 776 049 A (TAKAHASHI) 7 July 1998 (1998-07-07) column 5, line 3 -column 7, line 16 column 9, line 19 -column 10, line 19 figures 2A,B,9A,B ----	4
A	US 5 395 030 A (KURAMOTO ET AL.) 7 March 1995 (1995-03-07) column 5, line 56 -column 8, line 4 column 9, line 50 -column 13, line 45 column 26, line 43 -column 28, line 19 figures 1,6-14,58A-59C -----	7-10

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# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

IL 01/00843

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9100049	A	10-01-1991	DE 3921233 A1 AT 120628 T WO 9100049 A1 DE 59008855 D1 EP 0434793 A1 JP 4500768 T US 5166787 A	14-02-1991 15-04-1995 10-01-1991 11-05-1995 03-07-1991 13-02-1992 24-11-1992
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(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

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**Published:**

— without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

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WO 02/24058 A2

(54) Title: **MULTIPLE VIEW ENDOSCOPES**

(57) Abstract: The present invention relates to an endoscope comprising two or more optical channels that produce two or more distinct views. The endoscope of the invention is suitable for performing various surgical procedures, including fundoplications, stapling of the stomach for obesity management, bladder neck sling procedures for incontinence management, and other procedures that may benefit from having multiple interior views. Such treatments may be performed percutaneously, or by gaining access via natural body canals such as the esophagus or urethra.



## MULTIPLE VIEW ENDOSCOPES

### FIELD OF THE INVENTION

The invention applies to the field of endoscopy. More specifically the invention provides an endoscope with two or more optical channels that produce two or more distinct views.

### BACKGROUND OF THE INVENTION

GastroEsophageal Reflux Disease (GERD) is a common medical condition that affects approximately 44% of the US adult population to some degree. GERD symptoms are currently the most common reason that patients visit a gastroenterologist. The lower esophageal sphincter is a one-way valve that normally prevents stomach acids from entering the esophagus. Failure of the lower esophageal sphincter allows stomach acid to reflux upwards into the lower esophagus, typically after meals when the stomach is full and exerts greater pressure on the lower esophageal sphincter. Strong stomach acids will burn the delicate lining of the esophagus, causing "heartburn" or "acid indigestion" to occur. Heartburn is experienced as a burning or pressure sensation behind the breastbone, and can be accompanied by foul-tasting regurgitation into the back of the throat. Recurrent heartburn will damage the esophagus over time, and can lead to more serious conditions, including a cancer of the esophagus known as "Adenocarcinoma of the Gastroesophageal Cardia". Incidence of this particular type of cancer is rising in America and elsewhere.

Around 65 million Americans experience heartburn on a regular basis. The symptoms are uncomfortable, and most will seek some form of treatment. Options for treatment of acid reflux range from over-the-counter remedies and behavior modification (such as sitting up after meals or avoiding certain foods) to prescription drugs and surgery. Medications treat the symptoms of

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GERD, but do nothing to remedy the underlying cause. Drug treatments function by shielding the esophageal tissues for a period of time, stimulating their healing, or by suppressing stomach acid formation. All produce temporary effects and must be consumed on a regular basis. If the drugs are not taken, symptoms will very likely reoccur.

Surgical treatments are usually effective at eliminating the cause of acid reflux, and can be either open-incision or endoscopic (minimally invasive). The surgical procedure is designed to create a new functional lower esophageal sphincter or repair the damaged lower esophageal sphincter and thus prevent GERD. The most common surgical solution is to perform some variation of a procedure named "fundoplication". This surgery involves wrapping the fundus (upper body) of the stomach around the lower esophagus, which causes compression of the lower esophagus when the stomach is loaded with food. This effectively prevents acid reflux. The wrap of the fundus around the esophagus may be 360 degrees or less. More complete wraps are associated with inability or difficulty in belching or vomiting. Partial funduplications are therefore the preferred variation of this surgery. Funduplications may be performed in an open procedure, or using an endoscope with a percutaneous approach (inserting the endoscope through one or more incisions in the abdomen).

Very recently, trans-oral endoscopic procedures have been developed for surgical treatment of GERD. One procedure involves constricting the lower esophageal sphincter by suturing. Another procedure involves constricting the sphincter by injecting a material into the surrounding tissues to "fatten them up". Yet another procedure involves applying sufficient heat to surrounding tissues to stiffen them.

In copending International patent application PCT/IL01/00238 by the same applicant hereof, the description of which is incorporated herein by reference, there is described an endoscopic apparatus and procedure for performing

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partial funduplications that provides an alternative to all of the above-mentioned options for the treatment of GERD.

Endoscopy is a mature class of surgery that came into wide use after the invention of the Hopkins "rod-lens" relay system in the 1960s. Prior to this breakthrough, endoscopes provided very poor image quality coupled with an inability to provide and transmit adequate illumination and were not suitable for most surgical and diagnostic applications. An endoscope is an optical instrument used to visualize the interior of the human body, through natural or surgical openings. The advantages of endoscopic procedures include less trauma to the patient, shorter (or no) hospital stay, less pain, faster healing, and generally lower cost per procedure. The advantages of open surgery include greater ability for the physician to see and manipulate structures. The earliest endoscopes relied on the physician to directly view the interior surgical site by looking through the eyepiece of the endoscope. As video camera technology evolved, endoscopes could be coupled to a video camera indirectly through a coupling lens attached to the eyepiece, or directly by coupling the image to the sensor without use of an eyepiece at all. The use of video displays allows the entire operating team to view the surgical site, and the surgeon is not required to keep his eye at the endoscope ocular. The use of video also permits documentation (image storage) without the use of bulky and inconvenient photographic equipment.

Endoscopes currently exist in an array of different forms and are suitable for a wide variety of surgical procedures. Most endoscopes are designed to provide a broad view of the interior surgical site, but do not necessarily provide adequate visualization of the tools used with the endoscope. Even though endoscopes may be highly specialized for a particular procedure, they all contain the same basic component systems. An objective optical system captures a single image or view of the surgical area, a relay optical system carries the image from the distal to proximal end of the device, and an eyepiece or camera system (or both) are used to view the transmitted image.

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Light to illuminate the surgical scene is delivered via optical fibers or waveguides that are integral to the endoscope. The endoscope may also contain working channels or incorporate treatment options such as laser delivery. All of these parts are contained within an outer sheath that may be made from rigid or flexible materials. The endoscope itself may be rigid, semi-flexible, or flexible, and may have the ability to actively bend in one or more directions at its distal tip.

The objective of an endoscope may consist of glass or plastic lenses, diffractive or hybrid diffractive/refractive lenses, GRIN (graduated refractive index) lenses, prisms or mirrors. The image relay system may consist of a series of glass rods and lenses (a "rod lens" system), a series of lenses only, or fiberoptic image guides. The relay system may be bypassed in a video-only endoscope by placing the image sensor directly in the objective focal plane. The eyepiece typically consists of glass or plastic lenses. A video camera may be coupled to the eyepiece via a coupling lens, or may connect directly to the endoscope and view the image formed by the relay or objective system directly. A light source is coupled to the endoscope by a flexible fiberoptic cable in most cases, and is delivered by optical waveguides or fibers that may be glass or plastic. Some endoscopes provide viewing in stereo by incorporating more than one optical system at the proximal end to view the scene from two slightly offset perspectives. While these stereo endoscopes incorporate multiple image channels, they provide only one view of the surgical scene on an electronic display.

Endoscopes may be reusable or disposable, or may be split into one or more disposable and one or more reusable parts. Advantages of reusable endoscopes are that they are usually of much higher quality and have durability designed in. Disadvantages include degradation of the image quality after sterilization, which is performed using such methods as steam autoclave, ETO (ethylene oxide), glutaraldehyde, Steris (peracetic acid), Sterrad (hydrogen peroxide plasma), or other harsh chemicals and

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temperatures. The sterilization process degrades optical coatings, cements, and surfaces, and can also have deleterious effects on the mechanical parts. Another disadvantage of reusable endoscopes is their comparatively high initial cost. Disposable endoscopes do not suffer from repeated sterilization, and also reduce the possibility of cross-contamination from one surgical procedure to the next. Because they must be purchased in larger quantities and do not need to be as durable, initial costs are less than reusables (though per-use costs are typically higher). Endoscopes that are partly disposable and partly reusable are designed to maximize the advantages of each type of device while minimizing the disadvantages and cost per use.

### SUMMARY OF THE INVENTION

The present invention relates to an endoscope comprising two or more optical channels that produce two or more distinct views. The endoscope of the invention is suitable for performing various surgical procedures, including funduplications, stapling of the stomach for obesity management, bladder neck sling procedures for incontinence management, and other procedures that may benefit from having multiple interior views. Such treatments may be performed percutaneously, or by gaining access via natural body canals such as the esophagus or urethra.

Thus, in one aspect, the invention is directed to an endoscope comprising two or more separate optical channels that produce two or more distinct views, each of said optical channels consisting of an objective lens and a means of capturing or viewing the image; each channel optionally also including one or more of the following elements: a) an optical relay system; b) an ocular; and c) a coupling lens suitable to deliver the image acquired by said objective lens to an image sensor and display apparatus; wherein each objective lens is located at a different position along the length of the endoscope.

In another aspect, the invention is directed to a GERD endoscope comprising:

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- a) a sheath provided with a distal articulated section;
- b) stapler components distributed between a first location at the tip of said articulated section, and a second location along the length of said sheath, and which stapler components can be brought into a cooperative working positioned relationship by articulation of said articulating tip;
- c) a first objective lens located on said distal tip;
- d) a second objective lens located at said second location along the flexible sheath;
- e) a first optical channel to deliver the image acquired by said first objective lens to display apparatus coupled to said endoscope; and
- f) a second optical channel to deliver the image acquired by said second objective lens to display apparatus coupled to said endoscope.

In a further aspect, the invention is directed to a distal tip for a GERD endoscope comprising:

- a) a socket suitable to receive elements of a stapling device;
- b) at least one illumination channel; and
- c) at least one objective lens coupled to an optical relay system.

Each of the multiple views the endoscope produces has the following properties:

- The angle of the views may be between 0 to 180 degrees with respect to the mechanical axis of the endoscope.
- The field of view for each optical channel may be circular, with values up to 180 degrees or more.
- The field of view for each optical channel may be non-circular, such as square, rectangular, cylindrical, toroidal section, or other shape, or may be faceted, and may have an angular extent in each viewing axis of up to 180 degrees or more.
- For medical applications, the object viewed may be internal or external parts of human or animal bodies, parts of the endoscope, or surgical tools

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used in a procedure.

- For industrial applications, any object may be viewed, including parts of the endoscope and other tools used in the industrial procedure.

- The endoscope may operate in the visible, ultraviolet, infrared, or x-ray portions of the electromagnetic spectrum.

One or more of the following features may also be incorporated into the present invention:

- one or more image sensors to electronically acquire the views
- one or more displays to provide visual access to the electronically-acquired views
- one or more oculars to provide visual access to the views directly by the eye
- one or more of the image channels may provide stereoscopic imaging
- one or more portions that are disposable and one or more portions that are reusable
- one or more built-in functions (such as surgical staplers or laser delivery)
- one or more working channels for insertion of surgical instruments
- one or more illumination channels to provide adequate light for imaging or therapy such as photocuring or photoinitiation
- a rigid, semi-rigid, or flexible sheath, which may be disposable, resposable (multiple use disposable), or reusable
- active or passive articulation of the distal tip or an intermediate section in one or more axes

Each of the optical channels includes one or more objective lenses and may include one or more relay optical systems. An objective lens forms an image of a scene. A relay optical system carries an image from one location in space to a different location in space. In an endoscope, the relay system is used to carry the image formed by the objective from the objective image plane to the focal plane of the sensor or ocular that is used to provide visual access to the

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image. The objective optical system may include elements such as mirrors, prisms, glass and plastic lenses that may be spherical or aspheric, GRIN lenses, diffractive lenses, hybrid diffractive/refractive lenses, fresnel lenses, other optical elements, filters, apertures, mechanical spacers, and lens housings. The optics in the objective may have a fixed focal length, variable focal length ("continuous zoom"), or multiple focal lengths ("step zoom", or "power changer"). The relay optical system may consist of mirrors, prisms, glass and plastic lenses that may be spherical or aspheric, gradient index lenses, diffractive lenses, hybrid diffractive/refractive lenses, fresnel lenses, fiberoptic image guides, optically transparent rods (such as glass or plastic), apertures, mechanical spacers, and housings.

The endoscope provides viewing via ocular(s), image sensor(s), or both. Oculars are generally glass or plastic lenses that are specially designed to work in conjunction with the optics in the human eye. They may be used in conjunction with an image sensor by the addition of coupling optics in between the sensor and the ocular. The construction of the ocular is not limited to glass and plastic lenses, and may be comprised of more exotic optical elements such as diffractive, gradient index, fresnel, or other types of lenses. The optics in the ocular may have a fixed focal length, variable focal length ("continuous zoom"), or multiple focal lengths ("step zoom", or "power changer"). The endoscope may include a method of actively focusing the ocular, objective, or sensor, or these components may be fixed-focus.

Image sensors are typically optoelectronic devices that convert photons into electrical signals. In the visible spectrum, charge coupled devices (CCD) or complementary metal oxide semiconductor (CMOS) sensors are typically used. In the infrared spectrum, silicon, or lead salt sensors are typically used. In the ultraviolet spectrum, doped silicon sensors are typically used. In the X-ray spectrum, a wavelength-converting device such as a microchannel fiberoptic plate is used in conjunction with a CCD sensor. When an image sensor is used, a separate display is necessary to provide access to the image

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presented to the sensor.

If an electronic sensor(s) is used, at least one display will be used to present the image to one or more members of the surgical team. Display types include cathode ray tubes, flat panel displays, head-mounted displays, and volumetric displays. This display may be separate, or it may be incorporated into the endoscope itself. Multiple views may be imaged on a single display, or multiple displays may be utilized to present one or more of the multiple views individually. If more than one view is presented on the same display, the views may be tiled or otherwise arranged on the screen simultaneously, or there may be an option to toggle between the views, which are displayed one at a time. Other information may be presented on one or more of the displays. The other information displayed might include, but is not limited to, which view is active (in the case of toggled views), status of built-in surgical tools, or status of articulation (for endoscopes that include active articulation), status of sheath bend angle or shape (for flexible endoscopes), sensor readings, or vital signs (for endoscopes incorporating or in use with vital function monitoring devices).

Each of the multiple views may incorporate stereoscopic, monocular, or biocular viewing means. Monocular viewing presents a two-dimensional image to one eye through one ocular. Biocular viewing presents the same image to each eye, using two oculars, though this is still a two-dimensional image. Stereoscopic images are created by presenting each eye with a slightly offset view of the scene (call them "left" and "right" views). If the system is properly designed and fabricated, the human brain will "fuse" the left and right views. This simulates human vision, and produces enhanced depth perception. In an endoscope, stereoscopic images are produced by fabricating two optical channels inside the instrument. In a direct-view stereoscopic endoscope, these images are viewed by looking directly through dual oculars with the eyes (biocular vision). In an electronic stereoscopic endoscope, the two optical views are focused onto one or more electronic

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sensors and viewed via a stereoscopic or volumetric electronic display. A multiple view endoscope of the present invention provides distinct multiple views of one or more objects, and each view may or may not be stereoscopic. It is important to note that the "multiple views" of the present invention are not the same as the "slightly offset views" used by stereoscopic imaging systems. The viewing angles in the present invention are independent of each other, unlike those of stereoscopic imaging systems. The distinct views of the present invention are not intended to fuse.

The endoscope may be completely reusable, completely disposable, or be partly reusable and partly disposable. The disposable portion of the endoscope may contain all, part, or none of the optical elements. The disposable portion may contain all, part, or none of the built-in surgical tools. The disposable portion may or may not include the sensor or ocular.

The endoscope may incorporate built-in surgical tools. These tools may include, but are not limited to stapling mechanisms, drug delivery devices, suturing devices, cutting tools, laser delivery systems, cauterizing systems, spectroscopic instruments, irrigation systems, photoinitiation systems, photocuring systems, photodynamic therapy instruments, suction tools, sensors, electronic devices, and heating or cooling systems.

Alternatively, the endoscope may provide access for such tools by way of one or more working channels. In this case, the tools are not built into the endoscope, but are separate instruments that are guided to the surgical site by insertion into and through an unobstructed passageway within the endoscope.

The endoscope is provided with one or more illumination channels to provide the light necessary for imaging. Each view may have its own illumination source, or one illumination channel may provide adequate illumination for multiple views. Illumination channels include elements such as glass,

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plastic, or hollow optical fibers or optical waveguides. These are connected to an external light source via a liquid-filled or fiberoptic cable. These may employ mechanical adapters, which may incorporate lenses, fiberoptic tapers or other means to couple light efficiently from the external source into the endoscope illumination system. The endoscope may also provide light to the surgical site via built-in light sources, such as LEDs (light emitting diodes). Other sources of light may also be incorporated, either internally to the endoscope or externally, including transillumination of the surgical site, whereby light originating externally or in an adjacent body space reaches the site by passing through tissue.

The endoscope may be encased in a rigid, semi-rigid (maleable), or flexible outer sheath. The sheath may be disposable, resposable (multi-use disposable), or reusable.

Flexible endoscopes may have a built-in means of active articulation. Articulation may be employed to view different areas of the surgical site, to help guide the endoscope through convoluted paths inside the body, or to guide and place instruments during a surgical procedure. Articulation is controlled by the physician manipulating levers or other controls at the proximal end of the endoscope, or it may be controlled robotically or by other means.

All the above and other characteristics and advantages of the invention will be further understood through the following illustrative and non-limitative description of preferred embodiments thereof, with reference to the appended drawings.

#### **BRIEF DESCRIPTION OF THE DRAWINGS**

- Fig. 1 schematically shows a typical endoscope objective;
- Fig. 2A through Fig. 2E schematically show endoscope configurations

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with a single optical channel;

- Fig. 3A through Fig. 3C show different options for displaying the multiple views;
- Fig. 4A schematically shows the ocular types for monocular viewing;
- Fig. 4B schematically shows the ocular types for biocular viewing;
- Fig. 4C schematically shows the ocular types for stereoscopic viewing;
- Fig. 5A through Fig. 5D schematically show different configurations for the dual optical channels for dual views;
- Fig. 6 schematically shows a typical endoscope illumination system;
- Fig. 7 schematically shows an endoscope sheath with articulation;
- Fig. 8A schematically illustrates the fixed portion and the bending distal portion of the device of the invention;
- Fig. 8B schematically illustrates the bending of the endoscope of Fig. 5A through its fixed bending angle;
- Fig. 9A is a schematic view of an optical assembly showing dual optical channels;
- Fig. 9B is a cross-sectional view of the assembly of Fig. 8A looking from the distal end;
- Fig. 10 is a detailed view of the distal tip of the endoscope;
- Fig. 11 shows a display layout for a preferred embodiment of the invention;
- Figs. 12A, 12B, and 12C schematically illustrate the mechanical procedure involved in the fundoplication using a device according to the invention;
- Fig. 13 schematically illustrates the positioning of the device prior to stapling;
- Fig. 14A schematically shows the locking needles deployed from the anvil module and locked into the staple ejector module; and
- Fig. 14B schematically shows the situation after the staples have been fired and the locking needles withdrawn.

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## DETAILED DESCRIPTION OF THE INVENTION

Each of the multiple views of the invention is formed by an objective lens. Figure 1 shows the configuration of a typical endoscope objective (1) and illustrates some of its properties. The angle between the mechanical axis of the distal tip of the endoscope and the optical axis as it enters the endoscope objective ( $\alpha$ ) is the "angle of view". This angle of view refers only to the relationship between the optical and mechanical axes at the distal tip and does not take into account the variable direction of view provided by articulating (figure 7) the distal tip of the endoscope. Typical values for angle of view may range between 0 and 120 degrees. Non-zero angles of view are usually achieved by the use of prisms or mirrors in the objective optics. The field of view ( $\beta$ ) of the endoscope objective describes the angular extent in

object space that the lens can image. Field of view can be very narrow, approaching zero degrees, or may range up to 180 degrees. For example, a telescope will have a very small field of view with high magnification, while wide angle lenses (also known as "fish-eye" lenses due to the curved appearance of the image) have large fields of view with low magnification. The larger the field of view, the smaller the details will appear in the image. The image (3) of the object (2) can be located at or behind the last surface of the objective lens.

Looking at Fig. 2A through Fig. 2E, the image (3) formed by the objective lens (1) must be transmitted to a detector for viewing at the proximal end of the endoscope. An ocular (5) may be employed for direct viewing by the human eye (6), or the image may be formed on an optoelectronic sensor (7) (such as a CCD or CMOS sensor) that transforms the optical image into electronic signals. The electronic signals are then sent to a display, such as a video monitor, flat panel display, head-mounted display or liquid crystal display; where the image can be viewed by one or more people. Figures 2A to 2E show several configurations for each individual viewing channel of the

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present invention. Fig. 2A shows an objective lens (1) coupled to an ocular (5) by a relay optical system (4). A relay is an optical subsystem of the endoscope that transmits an image from the objective image plane to another location. Since an endoscope is usually much longer than the distance from the first surface of the objective to the objective image plane, relays are used to bring the image up to the proximal end of the endoscope, where it can be accessed by the viewing optics. Relays may consist of lenses or other optical imaging elements, or alternately a coherent (ordered) fiber optic image guide may be employed to transmit the image. Fiber optic image guides are usually used for flexible or semi-flexible endoscopes, while rigid endoscopes typically include a relay made up of a series of glass rods and lenses. Fig. 2B shows an objective lens (1) and relay (4) used in conjunction with an image sensor (7) connected optically via a coupling lens (8). The coupling lens images the relay's proximal image plane onto the surface of the sensor, where it is converted to an electronic signal and sent to the display for viewing. Fig. 2C shows an option in which an endoscope configured as shown in Fig. 1 is coupled to an image sensor (7) by a coupling lens (8) that clips or attaches onto the ocular (5) mount. Fig. 2D shows a configuration where the relay system is not used, and the image (3) is coupled directly from the objective image plane to the sensor surface (7) by a coupling lens (8). Fig. 2E is a similar option wherein the sensor surface may be placed directly in the objective image plane. This type of endoscope is sometimes referred to as a "chip-on-a-stick", and the sensor is imbedded in the distal end of the shaft instead of being located externally or at the proximal end.

If an image sensor is utilized, there are many options for displaying the multiple views provided by the invention. Single or multiple displays may be used, with single or multiple views on each display. Options include, but are not limited to those shown in Figures 3. In each scenario, the display may also convey other information such as status of built-in surgical tools, status of the articulation, or readouts from vital signs monitors for example. Fig. 3A illustrates having multiple views (1) and (2) arranged on a single display.

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The views may take on any shape and do not need to be of equal size. Fig. 3B shows multiple views (1) or (2) that are accessed on a single display by toggling between the views as needed. In another preferred embodiment of the present invention, a multiplicity of displays may provide visual access to a multiplicity of views. Each display may contain one or more views that can be accessed simultaneously, singly, or by toggling between views. Zero or more of the displays may also show status indicators or other information. As shown in Fig. 3C, one display is showing view (1) individually, a second display is toggling between views (2) and (6), and the third is presenting views (3), (4), and (5), simultaneously with status indicators (7).

Figures 4 show different ocular types for monocular, binocular, and stereoscopic viewing. It is important to note that while stereoscopic viewing may be provided for one or more of the multiple views, the optics used to provide a stereoscopic view do not present "multiple views" as defined with reference to the present invention. Key differences are that stereoscopic optical channels provide slightly offset images of the same object area; while in the present invention the optical channels that provide multiple views have substantial linear or angular offsets, or both, and do not provide images of the same object area. Fig. 4A illustrates the schematic for monocular viewing, which is typical of most endoscopes that include an ocular. One ocular (5) provides access to the image for a single eye (6), which can be either the left or the right eye. Biocular optics use splitting optics (9) to provide the same exact image to both eyes via two oculars, one for each eye as shown in Fig. 4B. Here the addition of the letters "L" and "R" designate left and right for both the oculars (5) and the eyes (6). The stereoscopic ocular arrangement shown in Figure 4C provides a slightly offset view of the image to each eye (6L and 6R) via two oculars (5L and 5R) that are offset. This simulates normal human vision where each eye captures a slightly offset view and allows for some depth perception.

According to a preferred embodiment of the present invention, each of the

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multiple views is formed by a unique objective lens (or pair of objective lenses for stereoscopic viewing). This means that at the distal end, there will be one optical channel per view (or two for stereoscopic viewing). These separate optical channels may continue through the entire endoscope as shown in Fig. 5A. This figure shows an endoscope having two views. Each view is carried to a separate image sensor (7) by its own unique optical channel. Since there are two views, there are two relays (4), two coupling lenses (8), and two sensors (7). In Fig. 5B, an alternative is shown where the endoscope has two views, that have unique objectives (1) and relay optical channels (4), and a single coupling lens (8) and image sensor (7) captures both views. The coupling lens and sensor may image the views together, or one at a time by switching between them actively or passively. A third alternative, shown in Figure 5C, has two unique objectives (1) that capture two views, with a single relay system (4), coupling lens (8), and sensor (7) providing access to the views. As in the system shown in Fig. 5B, the access may be simultaneous or one at a time. Fig. 5D illustrates an endoscope having two views, one of which provides stereoscopic imaging. The upper (non-stereo) view is carried by a single optical channel. The lower (stereo) view contains a set of two optical channels that are slightly offset at the distal end. The two optical channels present images of the object that originate from slightly different locations. This simulates normal human vision in which each eye views a scene from its own unique location and the brain integrates the differences in the image on each eye to formulate depth cues.

According to another preferred embodiment of the present invention, the endoscope provides illumination to the views via single or multiple illumination channels. Fig. 6 shows one such configuration where an endoscope with two views (as illustrated in Fig. 5A) has a separate illumination channel for each view. Light from an external source (20) is transmitted to the endoscope via liquid-filled or fiberoptic cable (21). Coupling optics (22) at the interface insure that the light is efficiently coupled into the internal illumination fibers (23). The illumination fibers are split

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into two channels (24 and 25) internally within the endoscope to provide light to each view individually.

The present invention utilizes a rigid, a semi-flexible, or a flexible sheath. If the endoscope is flexible, it may also be capable of articulation in one or more axes. Fig. 7 shows an example of 90-degree articulation in one axis on a flexible sheath.

As an illustrative, but not limitative, example of the invention, an endoscope with dual optical channels which permits carrying out a new surgical treatment for GastroEsophageal Reflux Disease (GERD) is now described.

The new procedure is an alternative to current treatment options and has been described in the above referenced international patent application (PCT/IL01/00238). This procedure is carried out using a surgical endoscopic device provided with at least one flexible portion, which is, in a preferred embodiment, an articulation section and a stapling device comprising a staple-firing portion and an anvil portion. According to preferred embodiments of the invention, the staple firing portion (sometimes called the cartridge) is located proximately to the proximal end of the articulation section and the anvil portion is located on the distal end or tip of the articulation section.

According to the preferred embodiment of the invention, the endoscope employs a two-way articulation system. In this case, completely bending the articulation section using a fixed radius of curvature brings the two portions of the stapler into alignment. Further, the parts of the stapling device are in correct working relationship when two alignment/locking pins that are stored in the anvil portion are extended and engage and lock into receptacles on the staple firing portion.

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The staple firing portion contains a staple cartridge containing one or a plurality of arrays of staples. Each array consists of one or a plurality of staples. The arrays of staples are fired by staple pushers actuated by cams actuatable by proximal means. The staple cartridge is indexable after the firing of each of the arrays of staples by the action of a proximal actuating device.

The endoscopic device for the GERD procedure should preferably comprise viewing means, typically a video camera. In a preferred embodiment of the invention, two separate optical channels are provided to provide two independent images, one from the area of the anvil at the distal tip and one from the area of the staple cartridge in the shaft of the endoscope. As will be apparent to the skilled person, it is usually necessary to provide illumination apparatus for viewing purposes. Additionally, other conventional endoscopic devices and accessories, such as water and/or air supply and/or suction, and/or ultrasound are provided.

Based on the above, an illustrative preferred embodiment of the present invention may, therefore, consist of the following elements:

- Multiple optical channels;
  - An optical channel at the distal tip to visualize insertion and stapling from the distal end (which also contains the stapler anvil);
  - A second optical channel located at the stapling backstop on the endoscope side wall to visualize staple penetration inside the esophagus from the "stapler side";
- A stapler module located intermediately along the endoscope shaft that may contain components of the optic system. This module is preferably disposable, but may be resposable or reusable;
- A stapler anvil module at the distal tip that may contain components of the optic system. This module is preferably disposable, but may be resposable or reusable;
- A robust, one-way articulating section e.g. capable of up to about 270° articulation to distend the fundus of the stomach and position the stapler;

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and

- A single display showing both views simultaneously, with optional status indicators for the endoscope articulation, stapling operation, or both.

Looking now at Fig. 8A, the distal portion of the device of the invention is schematically shown. This portion comprises a fixed, non-bending section, indicated at 2 (wherein is located the stapler ejector and the objective optics of the "stapler view" optical channel), and an articulating section 3, and the distal end 4 of length "l".

Articulating section 3 is similar in design to that of conventional endoscopes, but possesses several unique features. In order to simplify the alignment procedure and at the same time achieve maximum accuracy, a one-way articulation design has been chosen. This means that the articulating section is constrained to move in one direction only (i.e. the tip of the endoscope can only move from straight ahead to one side). Secondly, the device must be able to bend further than conventional endoscopes in order to carry out the required medical procedure. Finally the articulating section must be strong enough to provide a significant force against the tissues during fundus distension and stapling.

The fixed section, 2, contains the stapler cartridge. The stapler ejector has a side shooting design and requires an anvil, which is located on the end of the distal tip. Both the stapler cartridge and the anvil module are replaceable and fit into pockets on the shaft and distal tip. These pockets are labeled 1 and 1A respectively in Fig. 8A. The stapling elements at 1 and 1A, together, form the entire stapling assembly.

Articulation of the device is carried out in a conventional manner. Since the mechanism and operation of articulation activation are well known to the man of the art, they will not be discussed here for the sake of brevity.

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Fig. 8B shows the device of Fig. 8A in a fully bent position. The articulation section 3 has been bent through bending angle  $\phi$  using fixed radius of curvature  $r$ . The values of radius  $r$  and the length  $L$  (Fig. 8A) are determined by the fixed values  $l$  (length of the rigid distal tip) and  $y$  (the distance from the stapler cartridge to the junction of the fixed and bending portions of the distal end of the endoscope) in such a way that bending the device completely brings the two parts of the stapler assembly exactly into alignment.

Exact alignment is accomplished by deploying the locking pins to hold the endoscope in the position shown in Fig. 8B during the surgical procedure. In a preferred embodiment of the invention, locking pins are stored in the anvil section of the stapler. By using an activator (shown at 1 in Fig. 10), the physician can deploy the pins through the tissue of the fundus and esophagus walls and engage sockets in the staple ejector module. The method of deploying the locking pins is well known to the man of the art and will therefore not be discussed here for the sake of brevity.

Positioning markings 4 may be located on the device (as indicated in Fig. 8A), at the extremity outside the patient, to provide information on the length of device that has been introduced into the patient.

In order to see both sides of the staple as it is placed, and to assure proper joining of both fundus and esophageal tissues, a preferred embodiment of the invention employs the use of two optic channels (Fig. 9A). In this embodiment, an objective lens 1 captures the image from the tip of the scope (the "distal view"). A flexible fiberoptic image guide 2 carries the image about 12cm proximally where it is focused by a coupling lens 6 onto a CCD sensor 8. This view fills the main part of the video monitor (1 in Fig. 11), and is always displayed, since it is used during insertion, distention, and stapling. A "stapler view" (2 in Fig. 11) is simultaneously projected onto one corner of the CCD and thus appears in one corner of the monitor. This is a view from the

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endoscope shaft, looking sideways from the vicinity of the stapler, which is located at position 5 in Fig. 9A. The optical path of this image starts with an object perpendicular to the axis of the endoscope. The optical path travels through the stapler backstop 3 and clear portions of the stapler module and with the aid of right angle prism 4 and objective lens 7 an image is produced on the CCD 8. This view may only be activated during the stapling process. After stapling, the distal view shows the closed staple(s) from the stomach side, and the stapler view shows the staple(s) from the esophageal side. These multiple views provide confidence that each staple is properly placed before repositioning the instrument for the next shot.

The display may have the option of switching the second view off when it is not needed by controlling illumination to each view. The first view will typically be active continuously but also may be switched off.

Fig. 9B is a cross-sectional view of the optical system of Fig. 9A taken at A-A, looking toward the proximal end of the endoscope, showing how the various optical elements are arranged within the sheath of the endoscope 9.

Fig. 10 shows the distal tip of the endoscope. The anvil module of the stapler assembly goes into the socket 1. An actuator for locking pins that are contained in the anvil is shown at 2. A channel for suction or irrigation is shown at 5. The image channel is 4 and 3 represents illumination fibers.

The built-in surgical tools of the preferred embodiment of the invention consist of stapler and anvil modules and suction capability. A staple remover may also be incorporated. The stapler modules may be disposable or resposable items. The disposable portions of the endoscope may also contain parts of the optical channels, the outer sheath, a "bite block" for insertion of the scope orally, or other parts of the endoscope. The stapling module may place a single staple or a pattern of staples with each shot. The stapler contains one or more arrays of multiple staples that are advanced

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automatically after each shot.

The display may incorporate status indicators relating to the various functions of the endoscope. For example in Fig. 11, 3 shows the status of the articulation of the distal end, 4 shows the rotation of the endoscope around the long axis, and 5 shows the stapler status.

The partial fundoplication of the fundus of the stomach of a patient may be carried out with the endoscope of the invention by:

- a) moving the distal tip of the endoscopic device so as to engage the fundus of the patient and to displace it toward the lower part of the esophagus;
- b) bringing said stapling assembly into working positioned relationship by moving the distal portion of said endoscope through a fixed bending angle;
- c) determining when said two separate elements of said stapling assembly are aligned by viewing them through the viewing means of said endoscope and by the deployment of locking pins provided on the distal tip of said endoscope;
- d) ejecting a plurality of staples from said staple-ejecting device, thereby to connect the tissue between them;
- e) releasing the locking pins and straightening the distal bending portion of said endoscope; and
- f) rotating the endoscopic device relative to the axis of the esophagus and repeating steps (a) through (e) for as many times as needed to achieve the desired partial fundoplication.

The mechanical operation of the device involves the bending of the articulation section of the device so as to engage the fundus of the stomach with the distal tip, and to move it toward the lower esophagus. This is schematically illustrated in Figs. 12(A, B, and C). In Fig. 12A, two positions of the device are shown, a and a'. Position a' is the initial position after the device has been inserted through the mouth and esophagus of the patient to

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the desired position. Position a illustrates the beginning of bending of articulation section of the device, towards the fundus 6, the tip being indicated as 5.

In Fig. 12B, the bending of the device has proceeded to the stage in which the distal tip 5 has encountered the wall of the fundus 6 and started to pull it towards the lower region of the esophagus.

In Fig. 12C, the situation shown is that in which bending of the device has been completed, and the distal tip 5 has caused the fundus 6 to move from its original position to a position near the lower esophagus. In this position, the fundus is correctly positioned by tip 5 and it is possible to carry out the stapling together of the fundus and esophagus.

Fig. 13 is a more detailed view of the situation depicted in Fig. 12C. Here is schematically shown the alignment between the staple cartridge 1, mounted on the endoscope shaft 2 within the esophagus 7, and the anvil 1A mounted on the distal end 5 within the fundus 6.

In order to fasten the lower part of the fundus 6 (Fig. 13) to the lower part of the esophagus 7, by means of the stapling assembly, it is imperative that element 1 and element 1A be brought into the correct working positioned relationship, so that the staples, when ejected, perform their required task. Failure to bring the parts of the stapling assembly into the correct positioned relationship may be fatal, as it will result in the staple not being correctly positioned or folded, and in a high risk of damaging the tissue where the stapling has been performed.

As described above, the design of the device assures proper alignment. The surgeon is able to verify this alignment, as well as the proper distention of the fundus towards the esophagus, by using the visual means provided at the distal tip of the endoscope. Further, as the two parts of the stapler are

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pressed together, the tissue is pressed between them and it is possible to see through the tissue using the visual means provided on the cartridge side. Final alignment is accomplished by deploying the locking pins that are provided in the anvil section of the stapler. Figs. 14 shows the relevant part of the device and tissue. In Fig. 14A, the locking pins (collectively indicated at 11), that were stored in the anvil assembly 1A, have been deployed through the tissue of the fundus and esophagus walls and have engaged the sockets in the stapler cartridge 1. The locking pins not only perform the final alignment, but also increase the clamping force during the stapling which is now carried out.

Fig. 14B shows the situation after the stapling has been effected. Staples, (collectively indicated at 10), have engaged between the fundus and the esophagus, at the specific location on which it was operated. After carefully inspecting the staples the surgeon has retracted the locking pins. The surgeon next straightens the endoscope, and moves the device by rotating it to its next location. When the next location is reached, the bending/aligning procedure is repeated, and the stapling is effected again.

While embodiments of the invention have been described by way of illustration, it will be understood that the invention can be carried out by persons skilled in the art with many modifications, variations and adaptations, without departing from its spirit or exceeding the scope of the claims.

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## CLAIMS

1. An endoscope comprising two or more separate optical channels that produce two or more distinct views, each of said optical channels consisting of an objective lens and a means of capturing or viewing the image; each channel optionally also including one or more of the following elements: a) an optical relay system; b) an ocular; and c) a coupling lens suitable to deliver the image acquired by said objective lens to an image sensor and display apparatus; wherein each objective lens is located at a different position along the length of the endoscope.
2. An endoscope according to claim 1, in which each of said distinct multiple views may be formed by a single optical channel to produce a monocular view, or by multiple optical channels to produce a binocular or stereoscopic view.
3. An endoscope according to claim 1, in which the components of said optical channels and said display apparatus are chosen such that said endoscope can operate in either the visible, ultraviolet, infrared, or x-ray portions of the electromagnetic spectrum.
4. An endoscope according to claim 1, in which said objective lens, ocular, and coupling lens have either fixed focal length, multiple focal lengths, or variable focal lengths.
5. An endoscope according to claim 1, in which each of said distinct views is at an angle of between 0 and 180 degrees with respect to the mechanical axis of said endoscope.

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6. An endoscope according to claim 2, in which the field of view of each of each of said optical channels may be of any suitable shape, including, but not limited to circular and rectangular, and has an angular view of up to 180 degrees or more.
7. A GERD endoscope comprising:
  - a) a sheath provided with a distal articulated section;
  - b) stapler components distributed between a first location at the tip of said articulated section, and a second location along the length of said sheath, and which stapler components can be brought into a cooperative working positioned relationship by articulation of said articulating tip;
  - c) a first objective lens located on said distal tip;
  - d) a second objective lens located at said second location along the flexible sheath;
  - e) a first optical channel to deliver the image acquired by said first objective lens to display apparatus coupled to said endoscope; and
  - f) a second optical channel to deliver the image acquired by said second objective lens to display apparatus coupled to said endoscope.
8. A distal tip for a Gerd endoscope comprising:
  - a) a socket suitable to receive elements of a stapling device;
  - b) at least one illumination channel; and
  - c) at least one objective lens coupled to an optical relay system.
9. A distal tip according to claim 8, further comprising a suction and/or irrigation channel.
10. A distal tip according to claim 8, wherein the elements of the stapling device comprise an anvil.

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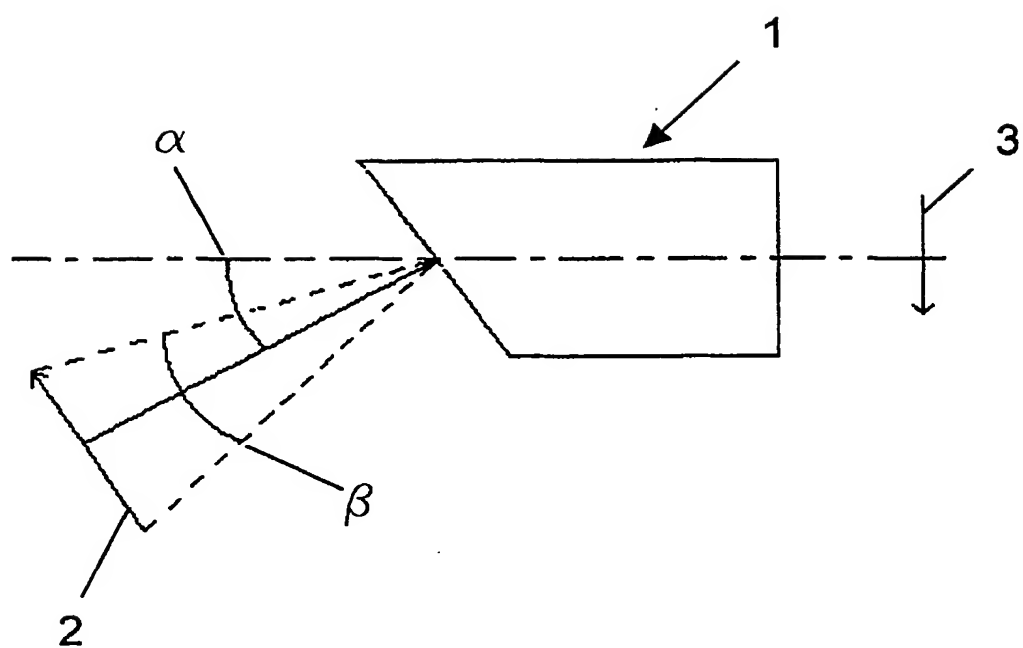


Fig. 1

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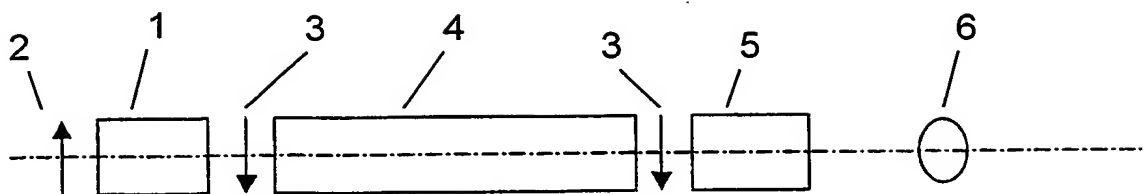


Fig. 2A

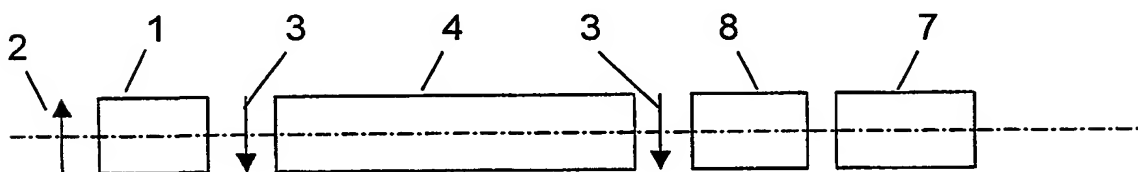


Fig. 2B

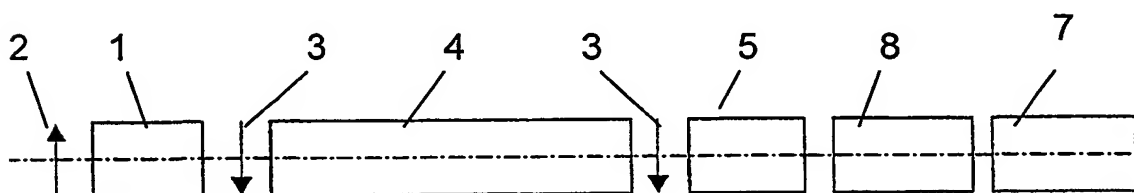


Fig. 2C

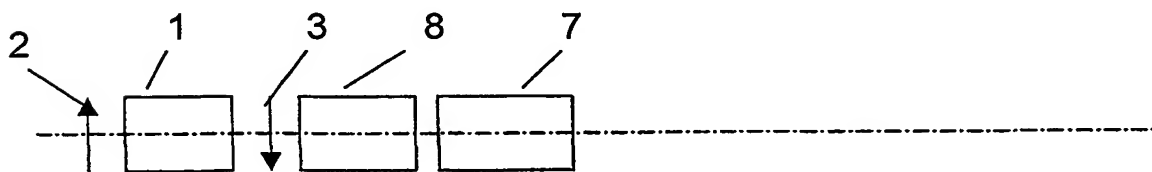


Fig. 2D

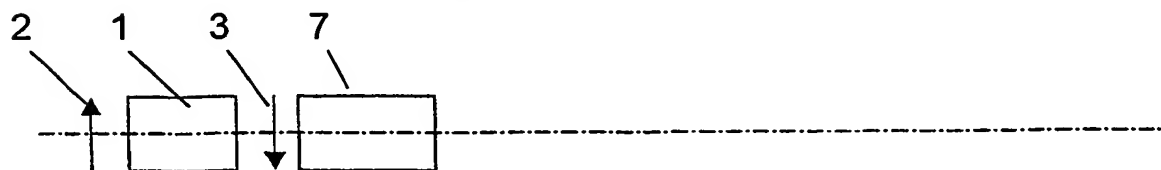


Fig. 2E

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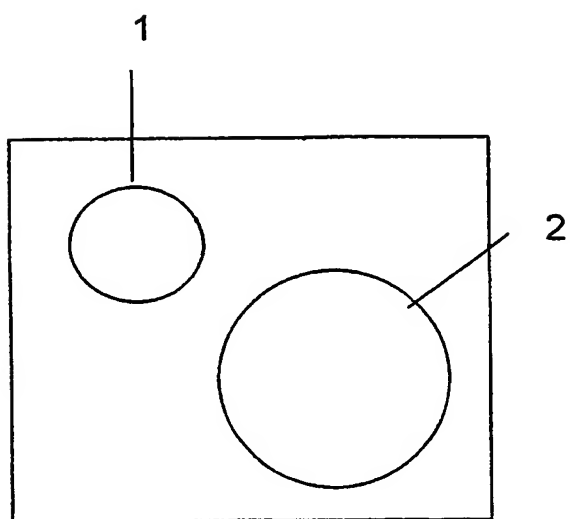


Fig. 3A

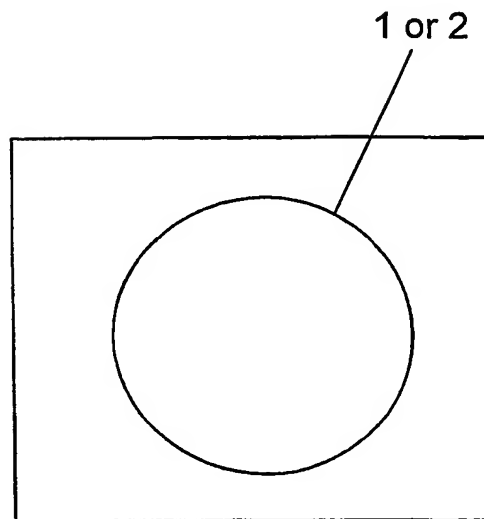


Fig. 3B

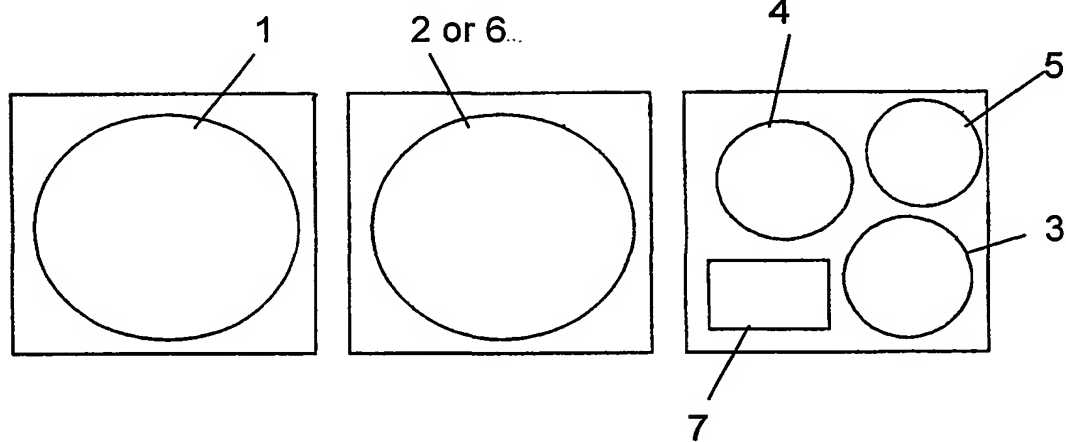


Fig. 3C

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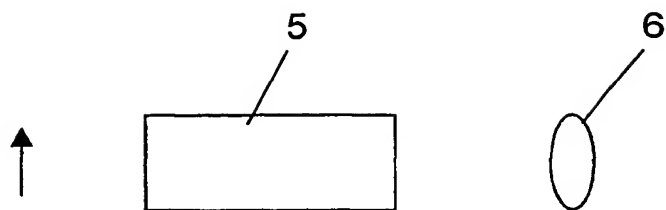


Fig. 4A

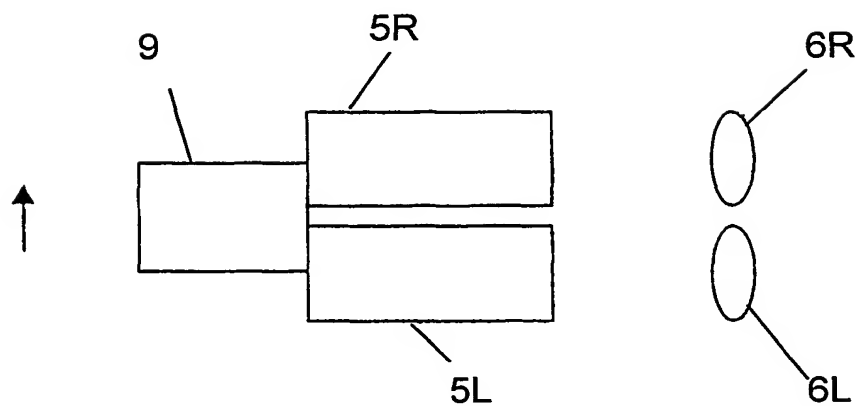


Fig. 4B

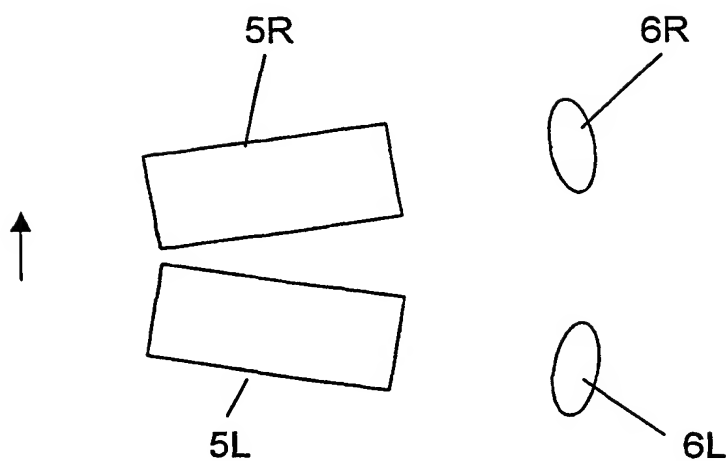


Fig. 4C

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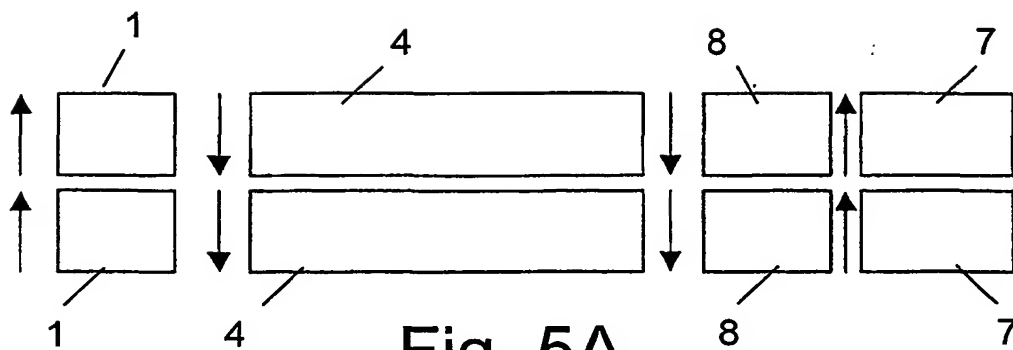


Fig. 5A

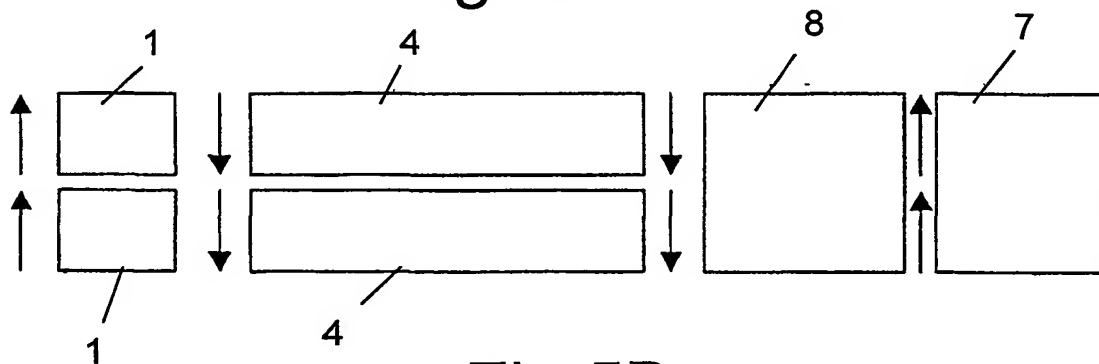


Fig. 5B

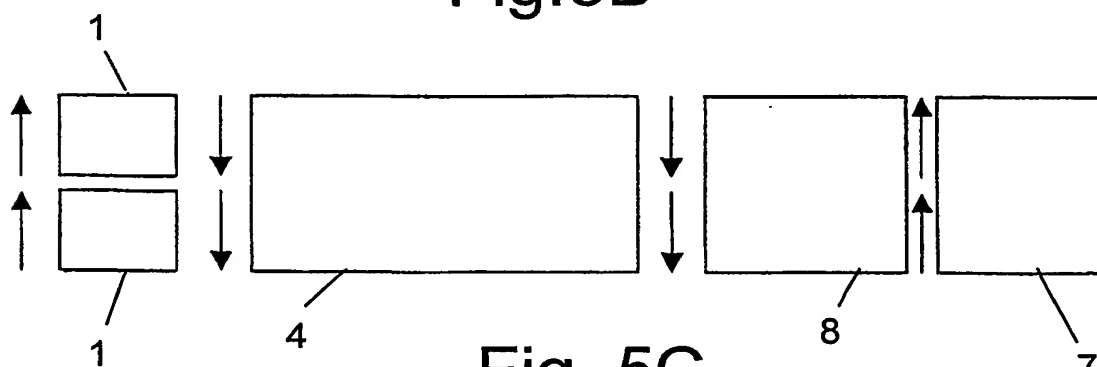


Fig. 5C

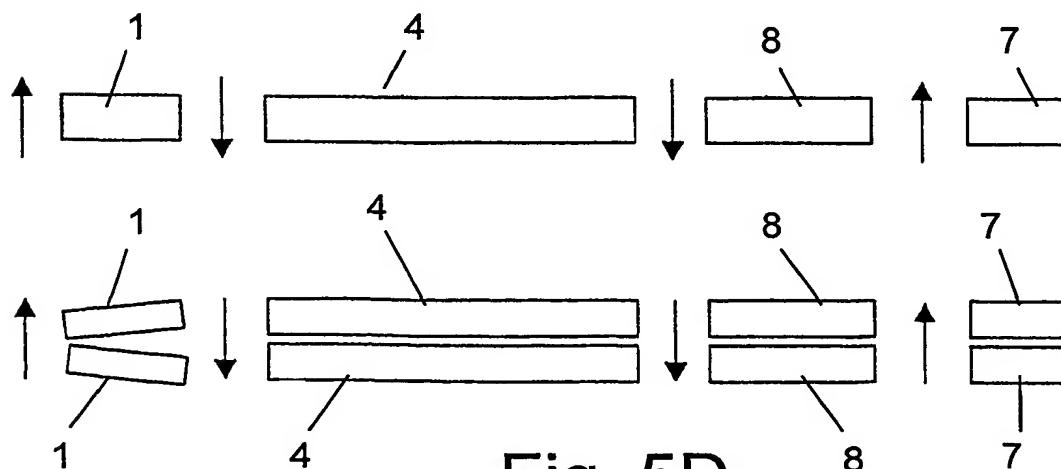


Fig. 5D

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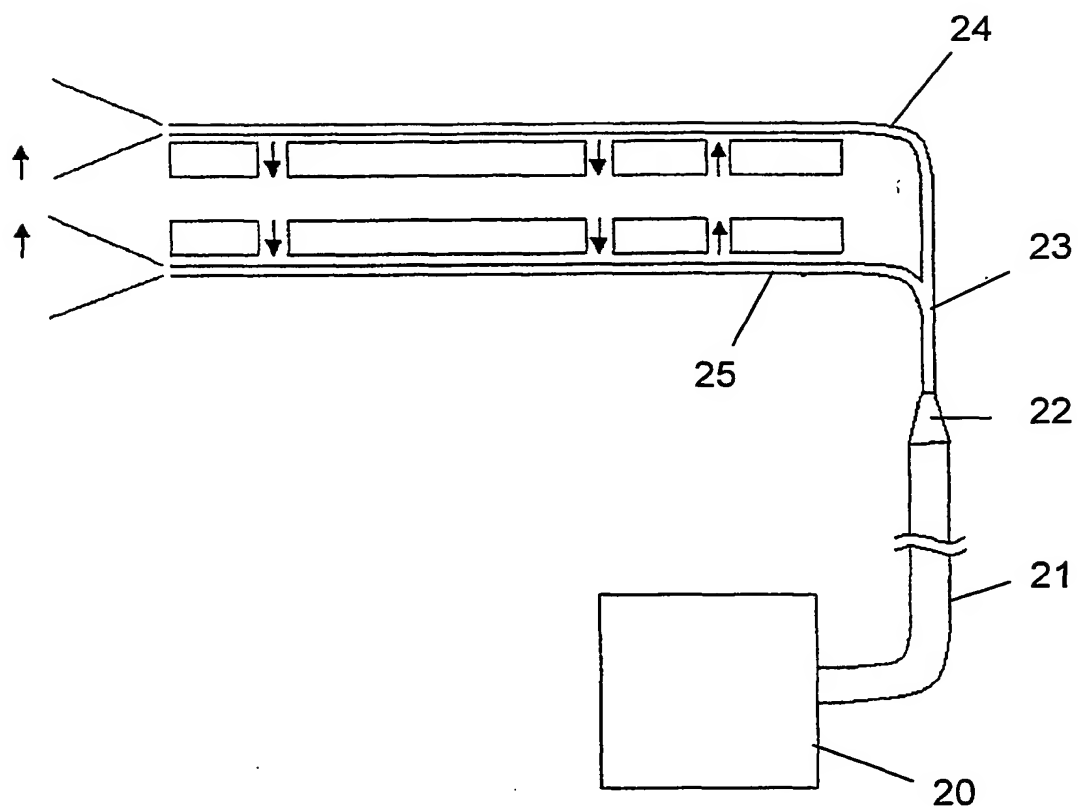


Fig. 6

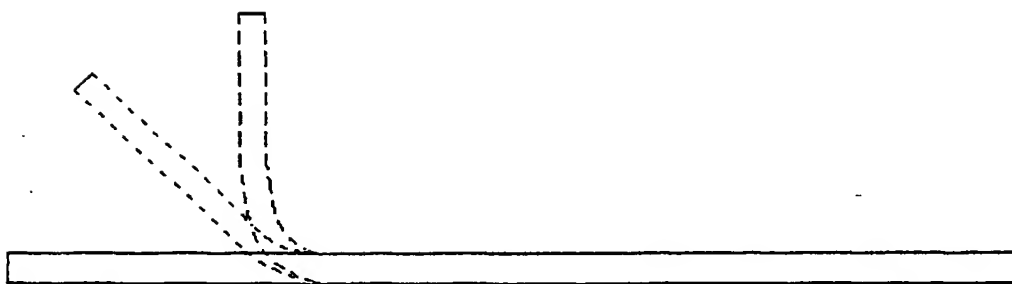


Fig. 7

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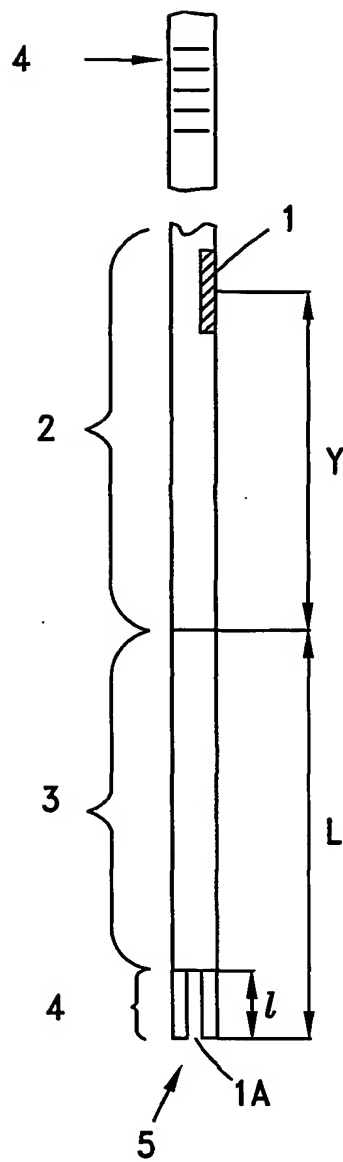


Fig. 8A

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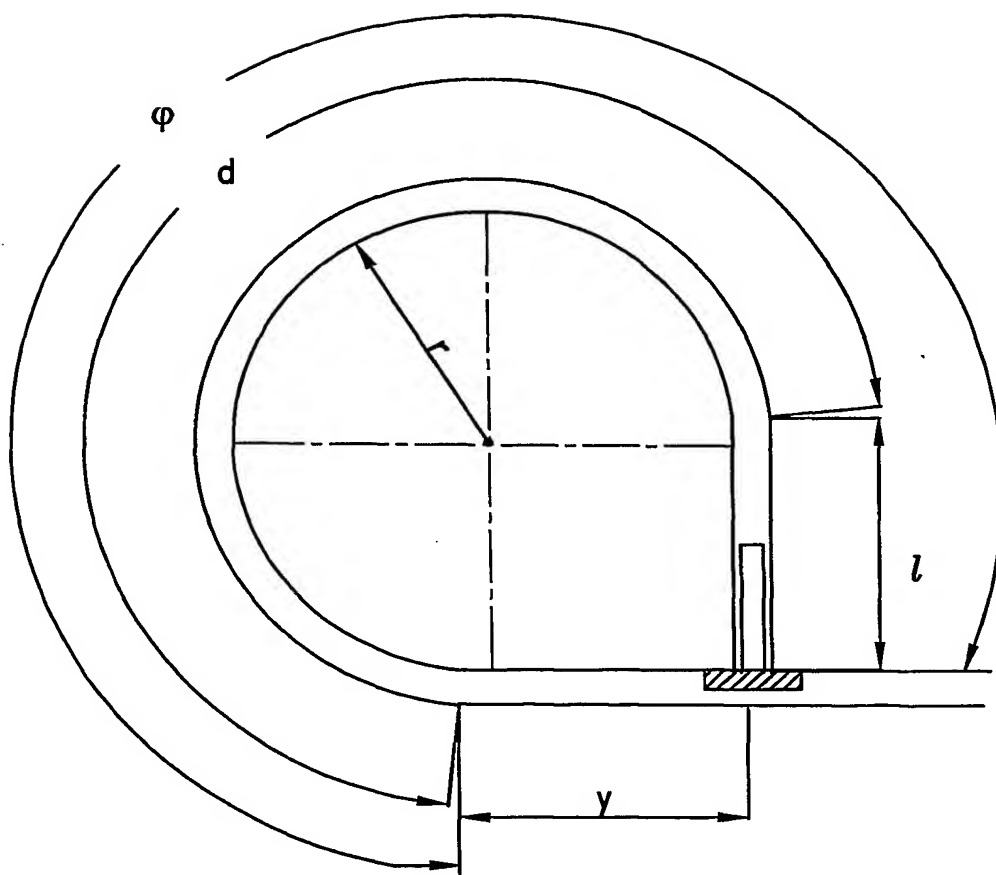


Fig. 8B

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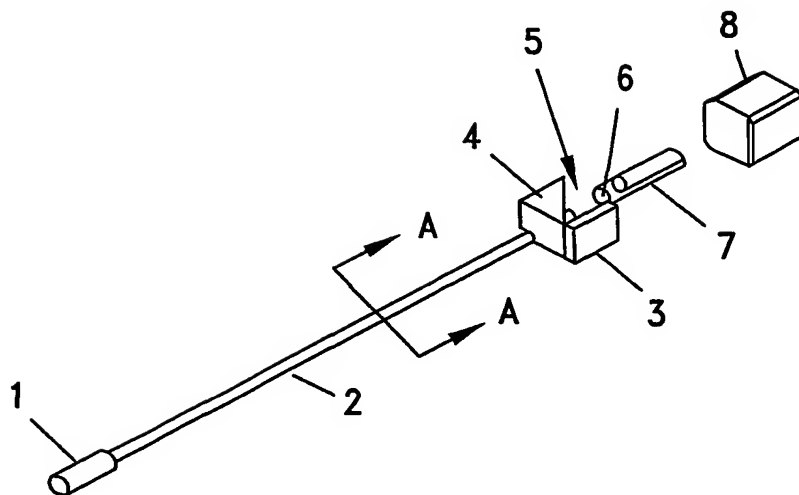


Fig. 9A

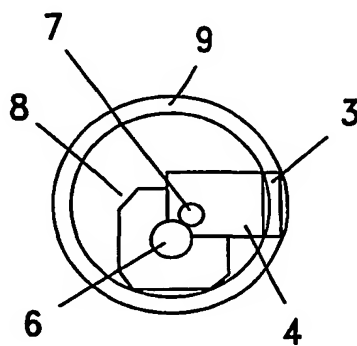


Fig. 9B

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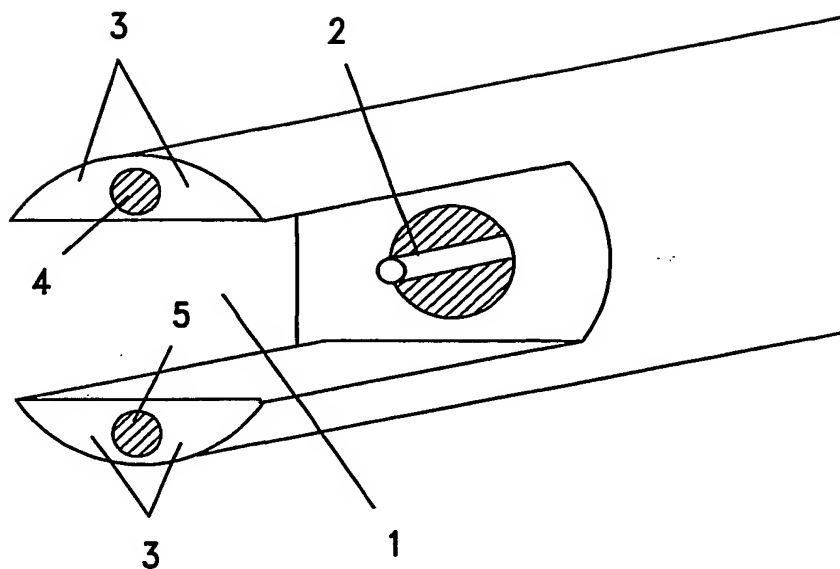


Fig. 10

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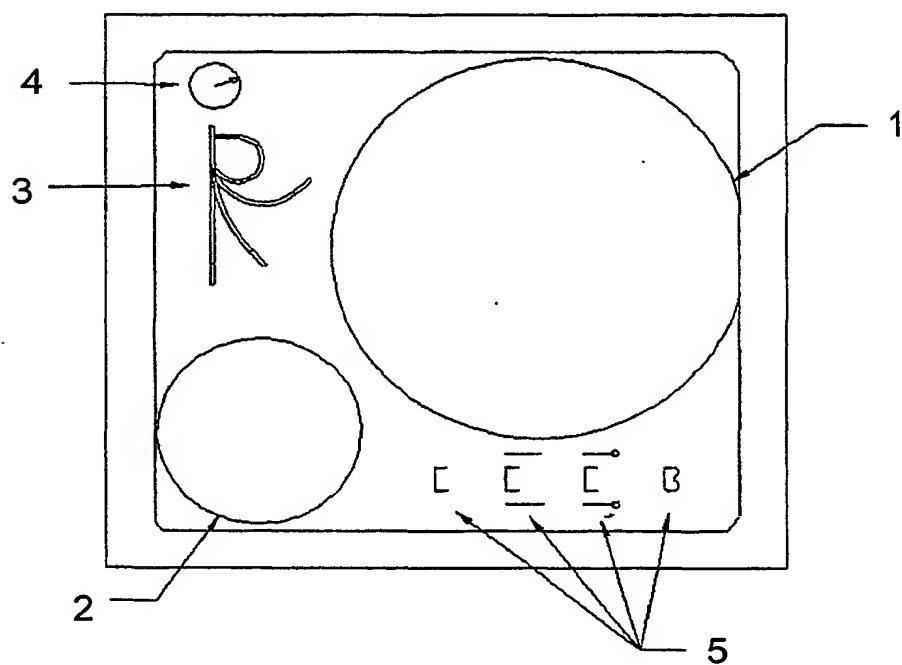


Fig. 11

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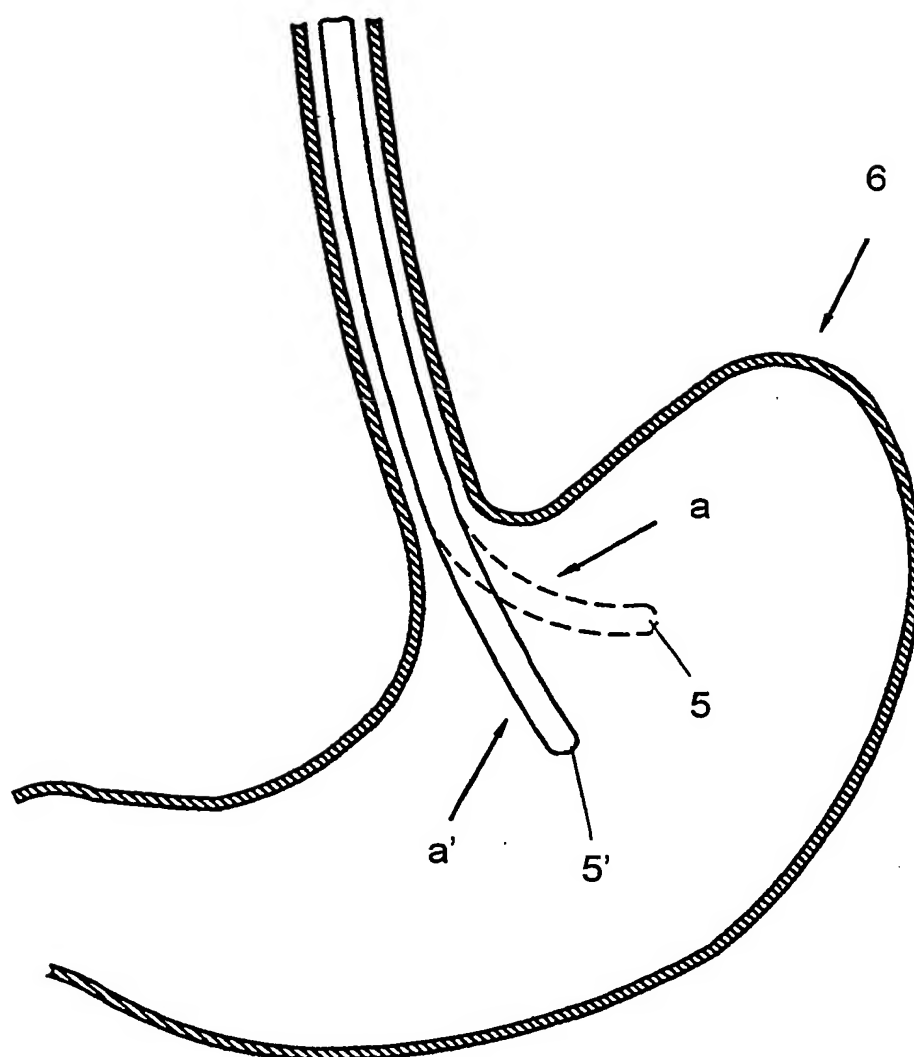


Fig. 12A

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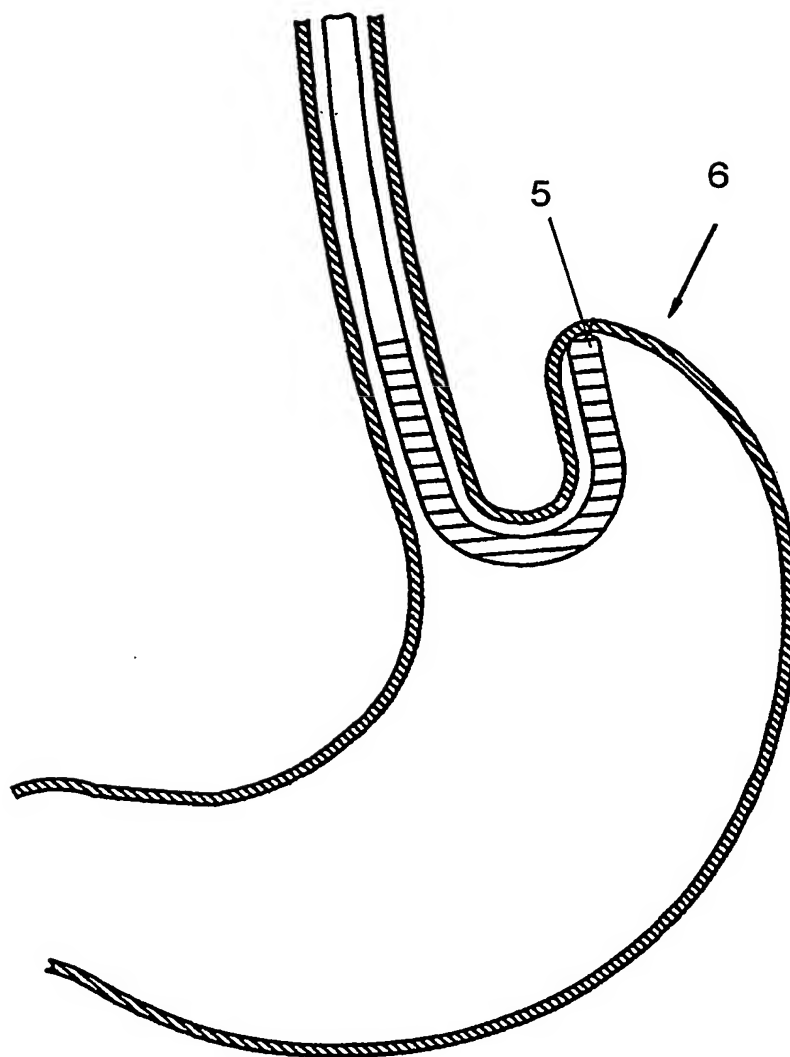


Fig. 12B

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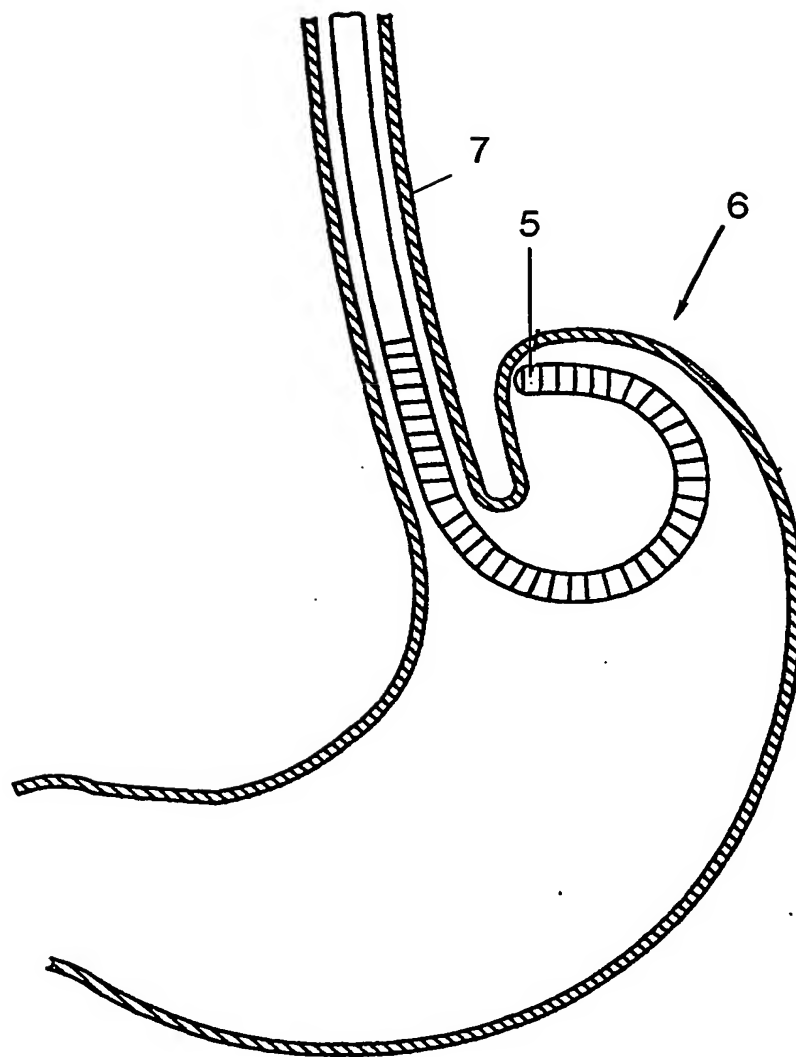


Fig. 12C

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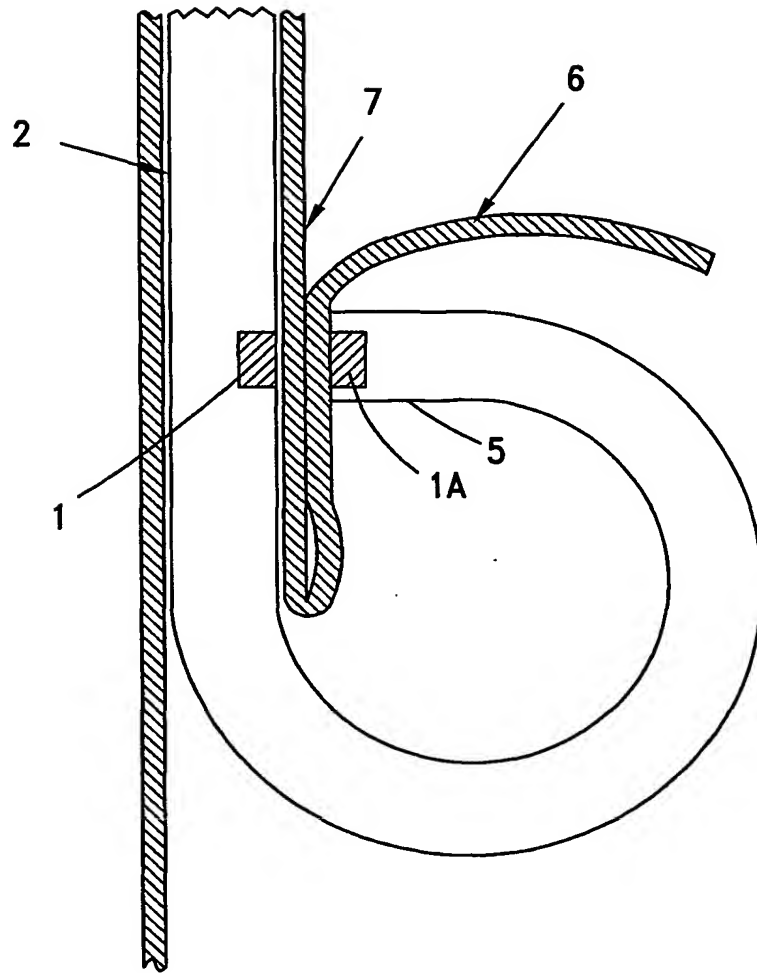


Fig. 13

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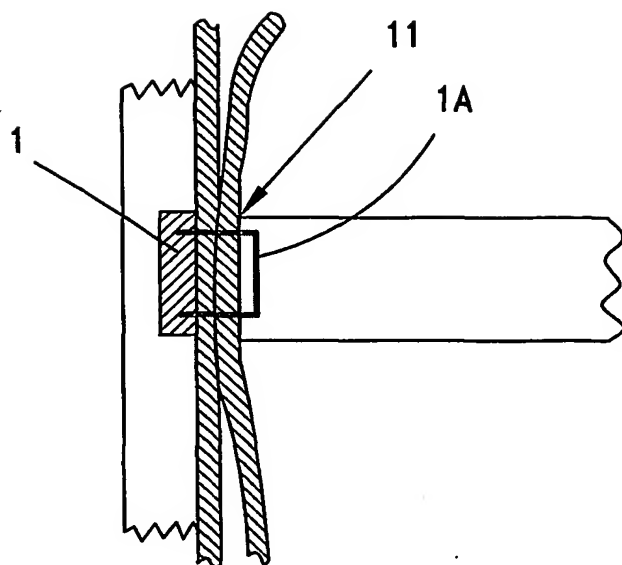


Fig. 14A

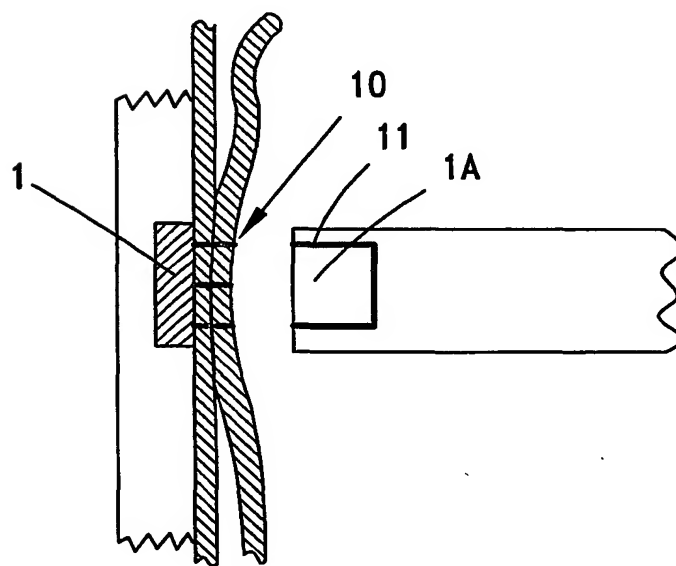


Fig. 14B

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(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
28 March 2002 (28.03.2002)

PCT

(10) International Publication Number  
**WO 02/024058 A3**

(51) International Patent Classification<sup>7</sup>: **A61B 1/00**

(21) International Application Number: PCT/IL01/00843

(22) International Filing Date:  
6 September 2001 (06.09.2001)

(25) Filing Language: English

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(30) Priority Data:  
138632 21 September 2000 (21.09.2000) IL

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(72) Inventors; and

(75) Inventors/Applicants (for US only): **SONNENSCHNEIN, Elazar** [IL/IL]; 32 Rahavat Yavne, 84800 Beersheva (IL). **SONNENSCHNEIN, Minelu** [IL/IL]; 12 Yelim Street, 85025 Meitar (IL). **CHINNOCK, Randal, B.** [US/US]; 53 McGilpin Road, Sturbridge, MA 01566 (US).

(74) Agents: **LUZZATTO, Kfir** et al.; Luzzatto & Luzzatto, P.O. Box 5352, 84152 Beer-Sheva (IL).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

**Published:**

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

(88) Date of publication of the international search report:  
22 August 2002

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: **MULTIPLE VIEW ENDOSCOPES**

(57) Abstract: The present invention relates to an endoscope comprising two or more optical channels that produce two or more distinct views. The endoscope of the invention is suitable for performing various surgical procedures, including funduplications, stapling of the stomach for obesity management, bladder neck sling procedures for incontinence management, and other procedures that may benefit from having multiple interior views. Such treatments may be performed percutaneously, or by gaining access via natural body canals such as the esophagus or urethra.

WO 02/024058 A3

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## INTERNATIONAL SEARCH REPORT

Int. Application No

PCT/1/00843

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61B1/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 91 00049 A (KARL STORZ GMBH & CO.) 10 January 1991 (1991-01-10) page 17, line 6 -page 23, line 14 figures 1-6C	1
A	---	2,4,5
X	DE 42 41 938 A (KARL STORZ GMBH & CO.) 16 June 1994 (1994-06-16) column 7, line 41 -column 9, line 33 figures	1
Y	---	7
Y	WO 00 53102 A (GAZAYERLI) 14 September 2000 (2000-09-14) page 7, line 20 -page 9, line 11 page 19, line 13 -page 23, line 3 figures 1-2B	7
A	---	8
	--- -/--	

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

## \* Special categories of cited documents:

\*A\* document defining the general state of the art which is not considered to be of particular relevance

\*E\* earlier document but published on or after the international filing date

\*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

\*O\* document referring to an oral disclosure, use, exhibition or other means

\*P\* document published prior to the international filing date but later than the priority date claimed

\*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

\*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*&amp;\* document member of the same patent family

Date of the actual completion of the international search

21 June 2002

Date of mailing of the international search report

- 8. 07. 2002

Name and mailing address of the ISA

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Authorized officer

Chen, A

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## INTERNATIONAL SEARCH REPORT

Int. Patent Application No.  
PC 01/00843

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3 889 662 A (MITSUI) 17 June 1975 (1975-06-17) column 4, line 4 - line 50 ----	1,6
A	WO 89 11252 A (UNIVERSITY OF CONNECTICUT) 30 November 1989 (1989-11-30) page 19, line 3 -page 24, line 11 figures 1,2,4,5 ----	1-3
A	US 5 776 049 A (TAKAHASHI) 7 July 1998 (1998-07-07) column 5, line 3 -column 7, line 16 column 9, line 19 -column 10, line 19 figures 2A,B,9A,B ----	4
A	US 5 395 030 A (KURAMOTO ET AL.) 7 March 1995 (1995-03-07) column 5, line 56 -column 8, line 4 column 9, line 50 -column 13, line 45 column 26, line 43 -column 28, line 19 figures 1,6-14,58A-59C -----	7-10

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# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/IL 01/00843

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☒ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☒ No protest accompanied the payment of additional search fees.

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**FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210**

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

**1. Claims: 1-6**

Endoscope comprising two or more separate optical channels, where each optical channel comprises an objective lens for focusing the image on to an image sensor chip. Each objective lens is located at a different position along the length of the endoscope.

**2. Claims: 7, 8-10**

Endoscope and surgical stapler combination.

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## INTERNATIONAL SEARCH REPORT

Information on patent family members

Int. Application No

PCT/JP01/00843

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9100049	A	10-01-1991	DE 3921233 A1	14-02-1991
			AT 120628 T	15-04-1995
			WO 9100049 A1	10-01-1991
			DE 59008855 D1	11-05-1995
			EP 0434793 A1	03-07-1991
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			CA 1316252 A1	13-04-1993
			EP 0416025 A1	13-03-1991
			JP 4500321 T	23-01-1992
			WO 8911252 A1	30-11-1989
			US 5109276 A	28-04-1992
			US 5200838 A	06-04-1993
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			JP 6194580 A	15-07-1994
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US 5395030	A	07-03-1995	JP 6047050 A	22-02-1994

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Patentamt

Zweigstelle  
in Den Haag

European  
Patent Office

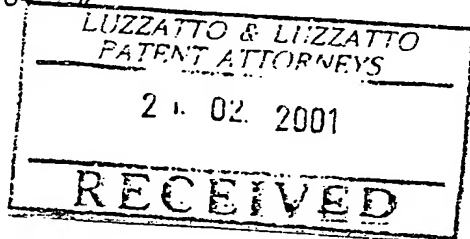
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Département à  
La Haye

1st Search Report

LUZZATTO & LUZZATTO  
P.O. Box 5352  
Beer-Sheva 84 152  
ISRAEL



Aktenzeichen/File No./No. du Dossier

RS 105958 IL

Datum/Date

Das Europäische Patentamt übermittelt hiermit den Standardrecherchenbericht zu dem unten bezeichneten Antrag; Kopien der im Recherchenbericht angeführten Schriften werden in der Anlage beigelegt.

The European Patent Office herewith transmits the Standard Search Report relating to the request indicated below; copies of the documents cited in the search report are enclosed.

L'Office Européen des Brevets à l'honneur de vous transmettre ci-joint le Rapport de Recherche concernant la demande désignée ci-dessous; des copies des documents cités sont jointes.

Zeichen und Datum des Antrages Applicant's reference and date Références et date de la demande	11777/00
Dokument, Gegenstand der Recherche Document subject of the search Objet de la recherche	ILA 13863200
Einreichungstag Filing date Date de dépôt	21/09/2000
Beanspruchte Priorität Priority claimed Priorité revendiquée	

OFFICE EUROPÉEN DES BREVETS  
Pour le Vice-Président,

R. de Best

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The Search Division considers that the present patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

1. Claims: 1-6

Endoscope comprising two or more separate optical channels, where each optical channel comprises an objective lens for focusing the image on to an image sensor chip. Each objective lens is located at a different position along the length of the endoscope.

2. Claims: 7, 8-10

Endoscope and surgical stapler combination.

The search has been limited to the invention first mentioned in the claims. A search for each of the other inventions can be carried out upon your request. For each additional invention a standard search fee will be due.

The prior art document is W09100049 (document D1).

This document describes an endoscope with two separate optical channels. In each channel, an objective lens receives the image and focuses it on to an image sensor chip. The objective lenses and image sensing chips are mounted in a distal swivelling mount, so that the two lenses may be positioned at two different longitudinal positions along the endoscope shaft (see p.22, 1.16 - p.23, 1.3 and figs 6a-c). The invention of claim 1 is, therefore, not new.

By comparison of the sub-claims, dependent on claim 1, and the disclosure D1, the features of claim 3 can be seen to make a contribution over D1 and can, therefore, be considered to be the "special technical features" of the first invention: optical components and display chosen such that the endoscope can operate in either visible, ultraviolet, infra-red or x-ray portions of the electromagnetic spectrum, solving the objectively determined problem of detecting and displaying images in a wide spectral range.

From comparison of the second invention of claims 7 and 8-10 and document D1, the "potential" STF of the second invention is a surgical stapler located in the endoscope, solving the objectively determined problem of combining an endoscope and a surgical stapler in one instrument.

The STFs of the two inventions are not the same, the objective problems related to the STFs are different and have no corresponding effect. Consequently, the requirements of Unity of Invention are not fulfilled.

One additional set of search fees is to be paid.

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DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
	LACK OF UNITY OF INVENTION see sheet B ---		
X	WO 91 00049 A (KARL STORZ GMBH & CO.) 10 January 1991 (1991-01-10) * page 17, line 6 - page 23, line 14 * * figures 1-6C *	1	
A	---	2,4,5	
X	DE 42 41 938 A (KARL STORZ GMBH & CO.) 16 June 1994 (1994-06-16) * column 7, line 41 - column 9, line 33 * * figures *	1	
A	---	1,6	
A	US 3 889 662 A (MITSUI) 17 June 1975 (1975-06-17) * column 4, line 4 - line 50 *	1,6	
A	---	1-3	
	WO 89 11252 A (UNIVERSITY OF CONNECTICUT) 30 November 1989 (1989-11-30) * page 19, line 3 - page 24, line 11 * * figures 1,2,4,5 *	1-3	TECHNICAL FIELDS SEARCHED (Int.CL.7)
A	---	4	A61B
	US 5 776 049 A (TAKAHASHI) 7 July 1998 (1998-07-07) * column 5, line 3 - column 7, line 16 * * column 9, line 19 - column 10, line 19 * * figures 2A,B,9A,B * -----	4	
-The present search report has been drawn up for all claims			
Date of completion of the search		Examiner	
9 February 2001		Chen, A	
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone			
Y : particularly relevant if combined with another document of the same category			
A : technological background			
O : non-written disclosure			
P : intermediate document			
T : theory or principle underlying the invention			
E : earlier patent document, but published on, or after the filing date			
D : document cited in the application			
L : document cited for other reasons			
-----			
& : member of the same patent family, corresponding document			

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## ANNEX TO THE STANDARD SEARCH REPORT NO.

RS 105958

This annex lists the patent family members relating to the patent documents cited in the above-mentioned search report.  
The members are as contained in the European Patent Office EDP file on  
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

09-02-2001

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9100049 A	10-01-1991	DE 3921233 A	14-02-1991
		AT 120628 T	15-04-1995
		DE 59008855 D	11-05-1995
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WO 8911252 A	30-11-1989	US 4938205 A	03-07-1990
		CA 1316252 A	13-04-1993
		EP 0416025 A	13-03-1991
		JP 4500321 T	23-01-1992
		US 5109276 A	28-04-1992
		US 5200838 A	06-04-1993
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US 5776049 A	07-07-1998	JP 6194580 A	15-07-1994
		US 5522789 A	04-06-1996

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# PATENT COOPERATION TREATY

# PCT

From the INTERNATIONAL SEARCHING AUTHORITY

To:  
LUZZATTO & LUZZATTO  
Attn. Luzzatto, Kfir  
P.O. Box 5352  
Beer-Sheva 84 152  
ISRAEL

## INVITATION TO PAY ADDITIONAL FEES

(PCT Article 17(3)(a) and Rule 40.1)

*Second Search Report*

Applicant's or agent's file reference <b>11777/WO/00</b>	Date of mailing (day/month/year) <b>26/02/2002</b>
International application No. <b>PCT/IL 01/00843</b>	International filing date (day/month/year) <b>06/09/2001</b>
Applicant <b>MEDIGUS LTD.</b>	

**1. This International Searching Authority**

- (i) considers that there are 2 (number of) inventions claimed in the international application covered by the claims indicated ~~XXX~~ on the extra sheet:

and it considers that the international application does not comply with the requirements of unity of invention (Rules 13.1, 13.2 and 13.3) for the reasons indicated ~~XXX~~ on the extra sheet:

- (ii) ☒ has carried out a partial international search (see Annex) ☐ will establish the international search report on those parts of the international application which relate to the invention first mentioned in claims Nos.:

**1-6**

- (iii) will establish the international search report on the other parts of the international application only if, and to the extent to which, additional fees are paid

**2. The applicant is hereby invited, within the time limit indicated above, to pay the amount indicated below:**

EUR 945,00 x 1 = EUR 945,00  
 Fee per additional invention      number of additional inventions      total amount of additional fees

Or, \_\_\_\_\_ x \_\_\_\_\_ = \_\_\_\_\_

The applicant is informed that, according to Rule 40.2(c), the payment of any additional fee may be made under protest, i.e., a reasoned statement to the effect that the international application complies with the requirement of unity of invention or that the amount of the required additional fee is excessive.

3. ☐ Claim(s) Nos. \_\_\_\_\_ have been found to be unsearchable under Article 17(2)(b) because of defects under Article 17(2)(a) and therefore have not been included with any invention.

Name and mailing address of the International Searching Authority



European Patent Office, P.B. 5818 Patentlaan 2  
NL-2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
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Authorized officer

**Johannes Van Brummelen**

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 PCT/ISA/206  
 2002-03-01

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**Annex to Form PCT/ISA/206  
COMMUNICATION RELATING TO THE RESULTS  
OF THE PARTIAL INTERNATIONAL SEARCH**

International Application No  
**PCT/IL 01/00843**

1. The present communication is an Annex to the invitation to pay additional fees (Form PCT/ISA/206). It shows the results of the international search established on the parts of the international application which relate to the invention first mentioned in claims Nos. 1-6
2. This communication is not the international search report which will be established according to Article 18 and Rule 43.
3. If the applicant does not pay any additional search fees, the information appearing in this communication will be considered as the result of the international search and will be included as such in the international search report.
4. If the applicant pays additional fees, the international search report will contain both the information appearing in this communication and the results of the international search on other parts of the international application for which such fees will have been paid.

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 91 00049 A (KARL STORZ GMBH & CO.) 10 January 1991 (1991-01-10) page 17, line 6 -page 23, line 14 figures 1-6C	1
A	----	2,4,5
X	DE 42 41 938 A (KARL STORZ GMBH & CO.) 16 June 1994 (1994-06-16) column 7, line 41 -column 9, line 33 figures	1
A	----	1,6
A	US 3 889 662 A (MITSUI) 17 June 1975 (1975-06-17) column 4, line 4 - line 50	1,6
A	----	1-3
A	WO 89 11252 A (UNIVERSITY OF CONNECTICUT) 30 November 1989 (1989-11-30) page 19, line 3 -page 24, line 11 figures 1,2,4,5	1-3
A	----	4
A	US 5 776 049 A (TAKAHASHI) 7 July 1998 (1998-07-07) column 5, line 3 -column 7, line 16 column 9, line 19 -column 10, line 19 figures 2A,B,9A,B	4

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

\* Special categories of cited documents :

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

- \*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- \*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- \*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- \*&\* document member of the same patent family

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# Patent Family Annex

Information on patent family members

International Application No

PCT/IL 01/00843

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
WO 9100049	A	10-01-1991	DE	3921233 A1	14-02-1991
			AT	120628 T	15-04-1995
			WO	9100049 A1	10-01-1991
			DE	59008855 D1	11-05-1995
			EP	0434793 A1	03-07-1991
			JP	4500768 T	13-02-1992
			US	5166787 A	24-11-1992
DE 4241938	A	16-06-1994	DE	4241938 A1	16-06-1994
			WO	9413190 A1	23-06-1994
			US	2002007110 A1	17-01-2002
US 3889662	A	17-06-1975	DE	2425827 A1	19-12-1974
WO 8911252	A	30-11-1989	US	4938205 A	03-07-1990
			CA	1316252 A1	13-04-1993
			EP	0416025 A1	13-03-1991
			JP	4500321 T	23-01-1992
			WO	8911252 A1	30-11-1989
			US	5109276 A	28-04-1992
			US	5200838 A	06-04-1993
			US	5172685 A	22-12-1992
US 5776049	A	07-07-1998	JP	3220538 B2	22-10-2001
			JP	6194580 A	15-07-1994
			US	5522789 A	04-06-1996

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This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-6

Endoscope comprising two or more separate optical channels, where each optical channel comprises an objective lens for focusing the image on to an image sensor chip. Each objective lens is located at a different position along the length of the endoscope.

2. Claims: 7, 8-10

Endoscope and surgical stapler combination.

The search has been limited to the invention first mentioned in the claims. A search for each of the other inventions can be carried out upon your request. For each additional invention a standard search fee will be due.

The prior art document is W09100049 (document D1).

This document describes an endoscope with two separate optical channels. In each channel, an objective lens receives the image and focuses it on to an image sensor chip. The objective lenses and image sensing chips are mounted in a distal swivelling mount, so that the two lenses may be positioned at two different longitudinal positions along the endoscope shaft (see p.22, 1.16 - p.23, 1.3 and figs 6a-c).

The invention of claim 1 is, therefore, not new.

By comparison of the sub-claims, dependent on claim 1, and the disclosure D1, the features of claim 3 can be seen to make a contribution over D1 and can, therefore, be considered to be the "special technical features" of the first invention: optical components and display chosen such that the endoscope can operate in either visible, ultraviolet, infra-red or x-ray portions of the electromagnetic spectrum, solving the objectively determined problem of detecting and displaying images in a wide spectral range.

From comparison of the second invention of claims 7 and 8-10 and document D1, the "potential" STF of the second invention is a surgical stapler located in the endoscope, solving the objectively determined problem of combining an endoscope and a surgical stapler in one instrument.

The STFs of the two inventions are not the same, the objective problems related to the STFs are different and have no corresponding effect. Consequently, the requirements of Unity of Invention are not fulfilled.

One additional set of search fees is to be paid.

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**INTERNATIONAL PRELIMINARY  
EXAMINATION REPORT**

International application No. PCT/IL01/00843

**I. Basis of the report**

1. With regard to the **elements** of the international application (*Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)*):

**Description, pages:**

1,2,5-13,16-18	as originally filed		
3,3A,4,14,15, 19-24	as received on	18/10/2002 with letter of	16/10/2002

**Claims, No.:**

1-9	with telefax of	01/12/2002
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**Drawings, sheets:**

1/16,2/16,4/16-6/16, 8/16	as originally filed		
3/16,7/16, 9/16-16/16	as received on	18/10/2002 with letter of	16/10/2002

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- ☐ the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☐ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

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## INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 11777/WO/00		FOR FURTHER ACTION See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
International application No. PCT/IL01/00843	International filing date (day/month/year) 06/09/2001	Priority date (day/month/year) 21/09/2000
International Patent Classification (IPC) or national classification and IPC A61B1/00		
Applicant MEDIGUS LTD. et al.		

- This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.
- This REPORT consists of a total of 5 sheets, including this cover sheet.  
  
☒ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

These annexes consist of a total of 23 sheets.



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- This report contains indications relating to the following items:

- ☒ Basis of the report
- ☐ Priority
- ☐ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- ☐ Lack of unity of invention
- ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- ☐ Certain documents cited
- ☐ Certain defects in the international application
- ☐ Certain observations on the international application

Date of submission of the demand 01/04/2002	Date of completion of this report 13.01.2003
Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Abraham, V Telephone No. +49 89 2399 7463 

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# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/IL01/00843

- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

- ☐ the description,      pages:  
☐ the claims,      Nos.:  
☐ the drawings,      sheets:

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

*(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)*

6. Additional observations, if necessary:

**V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement**

1. Statement

Novelty (N)	Yes: Claims 1-9
	No: Claims
Inventive step (IS)	Yes: Claims 1-9
	No: Claims
Industrial applicability (IA)	Yes: Claims 1-9
	No: Claims

2. Citations and explanations  
**see separate sheet**

**INTERNATIONAL PRELIMINARY  
EXAMINATION REPORT - SEPARATE SHEET**

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International application No. PCT/IL01/00843

Reference is made to the following documents:

- D1: WO 91 00049 A (KARL STORZ GMBH & CO.) 10 January 1991
- D2: DE 42 41 938 A (KARL STORZ GMBH & CO.) 16 June 1994
- D3: US-A-3 889 662 (MITSUI) 17 June 1975
- D4: US-A-5 395 030 (KURAMOTO ET AL.) 7 March 1995
- D5: WO 00 53102 A (GAZAYERLI) 14 September 2000

**V**

1. Document D4 which is at present considered to represent the most relevant prior art for the subject-matter of claim 1 discloses the following:

An endoscope (Fig. 22) comprising:  
a sheath provided with a distal articulated section (Fig. 22);  
stapler components (Fig. 22; 124,125,146);  
two or more separate optical channels (Fig. 22), each of said channels consisting of an objective lens (122,144) and a means for capturing or viewing the image (Fig. 22).

The problem to be solved by the additional features of claim 1 is to improve the handling of an endoscopic stapling device.

In D4 the stapler components and corresponding optical components are located on separate coaxially arranged sheaths which need to be operated separately. In document D5 the stapler components are located on the same sheath, however, the optical components are not located on this sheath, but a separate viewing device is provided. Documents D1-D3 define simple endoscopes without any stapler components. With the arrangement as defined in claim 1 only one sheath comprising both, the stapling and the optical components, needs to be operated. This combination of features is neither known nor rendered obvious by the available prior art. The requirements of Article 33(2)-(4) PCT are therefore met.

2. Claims 2-9 are dependent on claim 1 and therefore also meet the requirements of Article 33(2)-(4) PCT.

**INTERNATIONAL PRELIMINARY  
EXAMINATION REPORT - SEPARATE SHEET**

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International application No. PCT/IL01/00843

3. Claim 1 should have been drafted in the two part form in accordance with Rule 6.3(b) PCT, with those features known in combination from the prior art being placed in the preamble (Rule 6.3(b)(i) PCT) and with the remaining features being included in a characterising part (Rule 6.3(b)(ii) PCT).
4. The features of the claims should have been provided with reference signs placed in parentheses (Rule 6.2(b) PCT).
5. The passage of the description on page 24, last paragraph, referring to "the spirit of the invention" implies that the extent of protection may be expanded in a vague and not precisely defined way. Such formulations are not allowed (see PCT Guidelines C III 4.3a).

-3-

partial funduplications that provides an alternative to all of the above-mentioned options for the treatment of GERD.

So-called endoscopic surgical staplers of the prior art provide the cartridge and anvil as separate tools that are inserted into the cavity where they are to be used and are separately activated.

Typical of the prior art staplers are those disclosed in U.S. Patent No. 5,395,030. The staplers of the various embodiments disclosed in this patent are not integral to the endoscope. One component is contained at the distal face of the endoscope, but the second component is provided on a tool that is actuated through a working channel of the endoscope.

Endoscopy is a mature class of surgery that came into wide use after the invention of the Hopkins "rod-lens" relay system in the 1960s. Prior to this breakthrough, endoscopes provided very poor image quality coupled with an inability to provide and transmit adequate illumination and were not suitable for most surgical and diagnostic applications. An endoscope is an optical instrument used to visualize the interior of the human body, through natural or surgical openings. The advantages of endoscopic procedures include less trauma to the patient, shorter (or no) hospital stay, less pain, faster healing, and generally lower cost per procedure. The advantages of open surgery include greater ability for the physician to see and manipulate structures. The earliest endoscopes relied on the physician to directly view the interior surgical site by looking through the eyepiece of the endoscope. As video camera technology evolved, endoscopes could be coupled to a video camera indirectly through a coupling lens attached to the eyepiece, or directly by coupling the image to the sensor without use of an eyepiece at all. The use of video displays allows the entire operating team to view the surgical site, and the surgeon is not required to keep his eye at the endoscope ocular. The use of video also permits documentation (image storage) without the use of bulky and inconvenient photographic equipment.

-3A-

Endoscopes currently exist in an array of different forms and are suitable for a wide variety of surgical procedures. Most endoscopes are designed to provide a broad view of the interior surgical site, but do not necessarily provide adequate visualization of the tools used with the endoscope. Even though endoscopes may be highly specialized for a particular procedure, they all contain the same basic component systems. An objective optical system captures a single image or view of the surgical area, a relay optical system carries the image from the distal to proximal end of the device, and an eyepiece or camera system (or both) are used to view the transmitted image.

Light to illuminate the surgical scene is delivered via optical fibers or waveguides that are integral to the endoscope. The endoscope may also contain working channels or incorporate treatment options such as laser delivery. All of these parts are contained within an outer sheath that may be made from rigid or flexible materials. The endoscope itself may be rigid, semi-flexible, or flexible, and may have the ability to actively bend in one or more directions at its distal tip.

The objective of an endoscope may consist of glass or plastic lenses, diffractive or hybrid diffractive/refractive lenses, GRIN (graduated refractive index) lenses, prisms or mirrors. The image relay system may consist of a series of glass rods and lenses (a "rod lens" system), a series of lenses only, or fiberoptic image guides. The relay system may be bypassed in a video-only endoscope by placing the image sensor directly in the objective focal plane. The eyepiece typically consists of glass or plastic lenses. A video camera may be coupled to the eyepiece via a coupling lens, or may connect directly to the endoscope and view the image formed by the relay or objective system directly. A light source is coupled to the endoscope by a flexible fiberoptic cable in most cases, and is delivered by optical waveguides or fibers that may be glass or plastic. Some endoscopes provide viewing in stereo by incorporating more than one optical system at the

distal end to view the scene from two slightly offset perspectives. While these stereo endoscopes incorporate multiple image channels, they provide only one view of the surgical scene on an electronic display.

Examples of prior art endoscopes having two imaging systems at the distal tip are described in U.S. Patent No. 3,889,662 (corresponding to WO91/00049) and U.S. Patent Application No. 2002/0007110 (corresponding to DE 452 41 938). Other prior art solutions to the problem of providing two or more imaging systems involve providing at least one of the systems on a tool that projects from a working channel of the endoscope. Such systems are described in U.S. 5,166,787 and the above referenced U.S. 5,395,030.

None of the prior art endoscopes contain as an integral part dual optical systems that are separated along the longitudinal axis of the endoscope in order to allow observation of different areas within the cavity being examined and even of different parts of the endoscope itself.

Endoscopes may be reusable or disposable, or may be split into one or more disposable and one or more reusable parts. Advantages of reusable endoscopes are that they are usually of much higher quality and have durability designed in. Disadvantages include degradation of the image quality after sterilization, which is performed using such methods as steam autoclave, ETO (ethylene oxide), glutaraldehyde, Steris (peracetic acid), Sterrad (hydrogen peroxide plasma), or other harsh chemicals and



present invention. Fig. 2A shows an objective lens (1) coupled to an ocular (5) by a relay optical system (4). A relay is an optical subsystem of the endoscope that transmits an image from the objective image plane to another location. Since an endoscope is usually much longer than the distance from the first surface of the objective to the objective image plane, relays are used to bring the image up to the proximal end of the endoscope, where it can be accessed by the viewing optics. Relays may consist of lenses or other optical imaging elements, or alternately a coherent (ordered) fiber optic image guide may be employed to transmit the image. Fiber optic image guides are usually used for flexible or semi-flexible endoscopes, while rigid endoscopes typically include a relay made up of a series of glass rods and lenses. Fig. 2B shows an objective lens (1) and relay (4) used in conjunction with an image sensor (7) connected optically via a coupling lens (8). The coupling lens images the relay's proximal image plane onto the surface of the sensor, where it is converted to an electronic signal and sent to the display for viewing. Fig. 2C shows an option in which an endoscope configured as shown in Fig. 1 is coupled to an image sensor (7) by a coupling lens (8) that clips or attaches onto the ocular (5) mount. Fig. 2D shows a configuration where the relay system is not used, and the image (3) is coupled directly from the objective image plane to the sensor surface (7) by a coupling lens (8). Fig. 2E is a similar option wherein the sensor surface may be placed directly in the objective image plane. This type of endoscope is sometimes referred to as a "chip-on-a-stick", and the sensor is imbedded in the distal end of the shaft instead of being located externally or at the proximal end.

If an image sensor is utilized, there are many options for displaying the multiple views provided by the invention. Single or multiple displays may be used, with single or multiple views on each display. Options include, but are not limited to those shown in Figures 3. In each scenario, the display may also convey other information such as status of built-in surgical tools, status of the articulation, or readouts from vital signs monitors for example. Fig. 3A illustrates having multiple views (101) and (102) arranged on a single display.

The views may take on any shape and do not need to be of equal size. Fig. 3B shows multiple views (101) or (102) that are accessed on a single display by toggling between the views as needed. In another preferred embodiment of the present invention, a multiplicity of displays may provide visual access to a multiplicity of views. Each display may contain one or more views that can be accessed simultaneously, singly, or by toggling between views. Zero or more of the displays may also show status indicators or other information. As shown in Fig. 3C, one display is showing view (101) individually, a second display is toggling between views (102) and (106), and the third is presenting views (103), (104), and (105), simultaneously with status indicators (107).

Figures 4 show different ocular types for monocular, binocular, and stereoscopic viewing. It is important to note that while stereoscopic viewing may be provided for one or more of the multiple views, the optics used to provide a stereoscopic view do not present "multiple views" as defined with reference to the present invention. Key differences are that stereoscopic optical channels provide slightly offset images of the same object area; while in the present invention the optical channels that provide multiple views have substantial linear or angular offsets, or both, and do not provide images of the same object area. Fig. 4A illustrates the schematic for monocular viewing, which is typical of most endoscopes that include an ocular. One ocular (5) provides access to the image for a single eye (6), which can be either the left or the right eye. Biocular optics use splitting optics (9) to provide the same exact image to both eyes via two oculars, one for each eye as shown in Fig. 4B. Here the addition of the letters "L" and "R" designate left and right for both the oculars (5) and the eyes (6). The stereoscopic ocular arrangement shown in Figure 4C provides a slightly offset view of the image to each eye (6L and 6R) via two oculars (5L and 5R) that are offset. This simulates normal human vision where each eye captures a slightly offset view and allows for some depth perception.

According to a preferred embodiment of the present invention, each of the

and

A single display showing both views simultaneously, with optional status indicators for the endoscope articulation, stapling operation, or both.

Looking now at Fig. 8A, the distal portion of the device of the invention is schematically shown. This portion comprises a fixed, non-bending section, indicated at 202 (wherein is located the stapler ejector and the objective optics of the "stapler view" optical channel), and an articulating section 203, and the distal end 204 of length "L".

Articulating section 203 is similar in design to that of conventional endoscopes, but possesses several unique features. In order to simplify the alignment procedure and at the same time achieve maximum accuracy, a one-way articulation design has been chosen. This means that the articulating section is constrained to move in one direction only (i.e. the tip of the endoscope can only move from straight ahead to one side). Secondly, the device must be able to bend further than conventional endoscopes in order to carry out the required medical procedure. Finally the articulating section must be strong enough to provide a significant force against the tissues during fundus distension and stapling.

The fixed section, 202, contains the stapler cartridge. The stapler ejector has a side shooting design and requires an anvil, which is located on the end of the distal tip. Both the stapler cartridge and the anvil module are replaceable and fit into pockets on the shaft and distal tip. These pockets are labeled 201 and 201A respectively in Fig. 8A. The stapling elements at 201 and 20A, together, form the entire stapling assembly.

Articulation of the device is carried out in a conventional manner. Since the mechanism and operation of articulation activation are well known to the man of the art, they will not be discussed here for the sake of brevity.

Fig. 8B shows the device of Fig. 8A in a fully bent position. The articulation section 3 has been bent through bending angle  $\phi$  using fixed radius of curvature  $r$ . The values of radius  $r$  and the length  $L$  (Fig. 8A) are determined by the fixed values  $\ell$  (length of the rigid distal tip) and  $y$  (the distance from the stapler cartridge to the junction of the fixed and bending portions of the distal end of the endoscope) in such a way that bending the device completely brings the two parts of the stapler assembly exactly into alignment.

Exact alignment is accomplished by deploying the locking pins to hold the endoscope in the position shown in Fig. 8B during the surgical procedure. In a preferred embodiment of the invention, locking pins are stored in the anvil section of the stapler. By using an activator (shown at 402 in Fig. 10), the physician can deploy the pins through the tissue of the fundus and esophagus walls and engage sockets in the staple ejector module. The method of deploying the locking pins is well known to the man of the art and will therefore not be discussed here for the sake of brevity.

Positioning markings 204 may be located on the device (as indicated in Fig. 8A), at the extremity outside the patient, to provide information on the length of device that has been introduced into the patient.

In order to see both sides of the staple as it is placed, and to assure proper joining of both fundus and esophageal tissues, a preferred embodiment of the invention employs the use of two optic channels (Fig. 9A). In this embodiment, an objective lens 301 captures the image from the tip of the scope (the "distal view"). A flexible fiberoptic image guide 302 carries the image about 12cm proximally where it is focused by a coupling lens 306 onto a CCD sensor 308. This view fills the main part of the video monitor (501 in Fig. 11), and is always displayed, since it is used during insertion, distention, and stapling. A "stapler view" (502 in Fig. 11) is simultaneously projected onto one corner of the CCD and thus appears in one corner of the monitor. This is a view from the

endoscope shaft, looking sideways from the vicinity of the stapler, which is located at position 305 in Fig. 9A. The optical path of this image starts with an object perpendicular to the axis of the endoscope. The optical path travels through the stapler backstop 303 and clear portions of the stapler module and with the aid of right angle prism 304 and objective lens 307 an image is produced on the CCD 308. This view may only be activated during the stapling process. After stapling, the distal view shows the closed staple(s) from the stomach side, and the stapler view shows the staple(s) from the esophageal side. These multiple views provide confidence that each staple is properly placed before repositioning the instrument for the next shot.

The display may have the option of switching the second view off when it is not needed by controlling illumination to each view. The first view will typically be active continuously but also may be switched off.

Fig. 9B is a cross-sectional view of the optical system of Fig. 9A taken at A-A, looking toward the proximal end of the endoscope, showing how the various optical elements are arranged within the sheath of the endoscope 309.

Fig. 10 shows the distal tip of the endoscope. The anvil module of the stapler assembly goes into the socket 401. An actuator for locking pins that are contained in the anvil is shown at 402. A channel for suction or irrigation is shown at 405. The image channel is 404 and 403 represents illumination fibers.

The built-in surgical tools of the preferred embodiment of the invention consist of stapler and anvil modules and suction capability. A staple remover may also be incorporated. The stapler modules may be disposable or resposable items. The disposable portions of the endoscope may also contain parts of the optical channels, the outer sheath, a "bite block" for insertion of the scope orally, or other parts of the endoscope. The stapling module may place a single staple or a pattern of staples with each shot. The stapler contains one or more arrays of multiple staples that are advanced

automatically after each shot.

The display may incorporate status indicators relating to the various functions of the endoscope. For example in Fig. 11, 503 shows the status of the articulation of the distal end, 504 shows the rotation of the endoscope around the long axis, and 505 shows the stapler status.

The partial fundoplication of the fundus of the stomach of a patient may be carried out with the endoscope of the invention by:

- a) moving the distal tip of the endoscopic device so as to engage the fundus of the patient and to displace it toward the lower part of the esophagus;
- b) bringing said stapling assembly into working positioned relationship by moving the distal portion of said endoscope through a fixed bending angle;
- c) determining when said two separate elements of said stapling assembly are aligned by viewing them through the viewing means of said endoscope and by the deployment of locking pins provided on the distal tip of said endoscope;
- d) ejecting a plurality of staples from said staple-ejecting device, thereby to connect the tissue between them;
- e) releasing the locking pins and straightening the distal bending portion of said endoscope; and
- f) rotating the endoscopic device relative to the axis of the esophagus and repeating steps (a) through (e) for as many times as needed to achieve the desired partial fundoplication.

The mechanical operation of the device involves the bending of the articulation section of the device so as to engage the fundus of the stomach with the distal tip, and to move it toward the lower esophagus. This is schematically illustrated in Figs. 12(A, B, and C). In Fig. 12A, two positions of the device are shown, a and a'. Position a' is the initial position after the device has been inserted through the mouth and esophagus of the patient to

the desired position. Position a illustrates the beginning of bending of articulation section of the device, towards the fundus 606, the tip being indicated as 205.

In Fig. 12B, the bending of the device has proceeded to the stage in which the distal tip 205 has encountered the wall of the fundus 606 and started to pull it towards the lower region of the esophagus.

In Fig. 12C, the situation shown is that in which bending of the device has been completed, and the distal tip 205 has caused the fundus 606 to move from its original position to a position near the lower esophagus. In this position, the fundus is correctly positioned by tip 205 and it is possible to carry out the stapling together of the fundus and esophagus.

Fig. 13 is a more detailed view of the situation depicted in Fig. 12C. Here is schematically shown the alignment between the staple cartridge 201, mounted on the endoscope shaft 202 within the esophagus 607, and the anvil 201A mounted on the distal end 205 within the fundus 606.

In order to fasten the lower part of the fundus 606 (Fig. 13) to the lower part of the esophagus 607, by means of the stapling assembly, it is imperative that element 201 and element 201A be brought into the correct working positioned relationship, so that the staples, when ejected, perform their required task. Failure to bring the parts of the stapling assembly into the correct positioned relationship may be fatal, as it will result in the staple not being correctly positioned or folded, and in a high risk of damaging the tissue where the stapling has been performed.

As described above, the design of the device assures proper alignment. The surgeon is able to verify this alignment, as well as the proper distention of the fundus towards the esophagus, by using the visual means provided at the distal tip of the endoscope. Further, as the two parts of the stapler are

pressed together, the tissue is pressed between them and it is possible to see through the tissue using the visual means provided on the cartridge side. Final alignment is accomplished by deploying the locking pins that are provided in the anvil section of the stapler. Figs. 14 shows the relevant part of the device and tissue. In Fig. 14A, the locking pins (collectively indicated at 11), that were stored in the anvil assembly 201A, have been deployed through the tissue of the fundus and esophagus walls and have engaged the sockets in the stapler cartridge 201. The locking pins not only perform the final alignment, but also increase the clamping force during the stapling which is now carried out.

Fig. 14B shows the situation after the stapling has been effected. Staples, (collectively indicated at 10), have engaged between the fundus and the esophagus, at the specific location on which it was operated. After carefully inspecting the staples the surgeon has retracted the locking pins. The surgeon next straightens the endoscope, and moves the device by rotating it to its next location. When the next location is reached, the bending/aligning procedure is repeated, and the stapling is effected again.

While embodiments of the invention have been described by way of illustration, it will be understood that the invention can be carried out by persons skilled in the art with many modifications, variations and adaptations, without departing from its spirit or exceeding the scope of the claims.



CLAIMS

1. An endoscope comprising a sheath provided with a distal articulated section, stapler components, and two or more separate optical channels that produce two or more distinct views, each of said optical channels consisting of an objective lens and a means of capturing or viewing the image, each channel optionally also including one or more of the following elements: a) an optical relay system; b) an ocular; and c) a coupling lens suitable to deliver the image acquired by said objective lens to an image sensor and display apparatus;

wherein said stapler components are distributed between a first location at the distal tip of said articulated section, and a second location, located adjacent to or proximally of the proximal end of said articulation section along the length of said sheath, and which stapler components can be brought into a cooperative working positioned relationship by articulation of said articulation section; and

wherein, a first of said objective lenses is located on said distal tip adjacent to the first of said stapler components and a second of said objective lenses is located adjacent to the second of said stapler components at said second location.

2. An endoscope according to claim 1, in which each of said distinct multiple views may be formed by a single optical channel to produce a monocular view, or by multiple optical channels to produce a binocular or stereoscopic view.

3. An endoscope according to claim 1, in which the components of said optical channels and said display apparatus are chosen such that said endoscope can operate in either the visible, ultraviolet, infrared, or x-ray portions of the electromagnetic spectrum.

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4. An endoscope according to claim 1, in which said objective lens, ocular, and coupling lens have either fixed focal length, multiple focal lengths, or variable focal lengths.
5. An endoscope according to claim 1, in which each of said distinct views is at an angle of between 0 and 180 degrees with respect to the mechanical axis of said endoscope.
6. An endoscope according to claim 1, in which the field of view of each of each of said optical channels may be of any suitable shape, including, but not limited to circular and rectangular, and has an angular view of up to 180 degrees or more.
7. A distal tip for an endoscope according to claim 1, further comprising:
  - a) a socket suitable to receive elements of a stapling device;
  - b) at least one illumination channel; and
  - c) at least one objective lens coupled to an optical relay system.
8. A distal tip according to claim 7, further comprising a suction and/or irrigation channel.
9. A distal tip according to claim 7, wherein the elements of the stapling device comprise an anvil.

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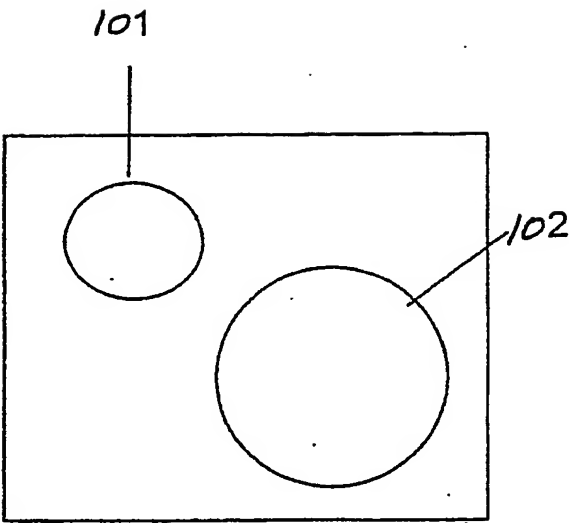


Fig. 3A

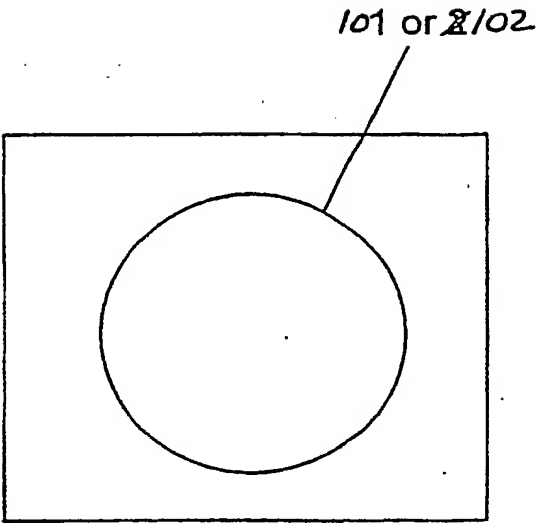


Fig. 3B

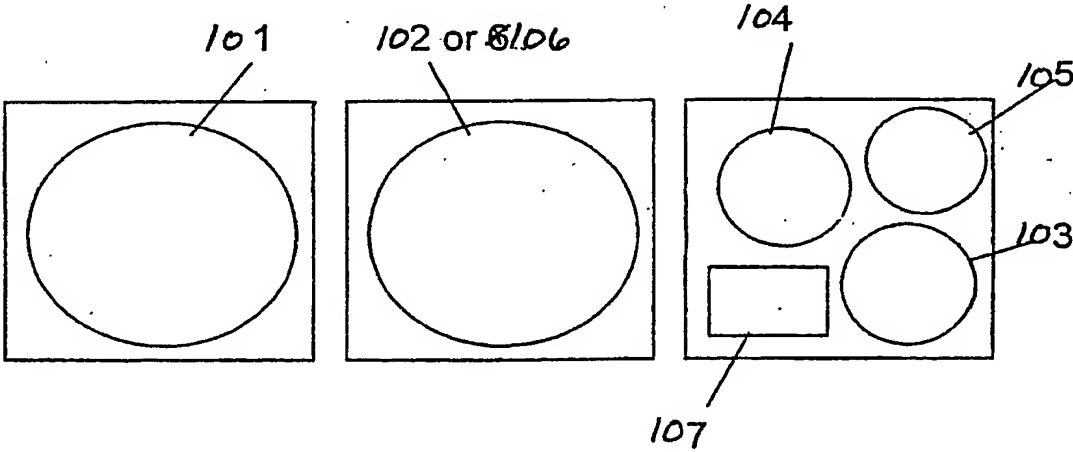


Fig. 3C

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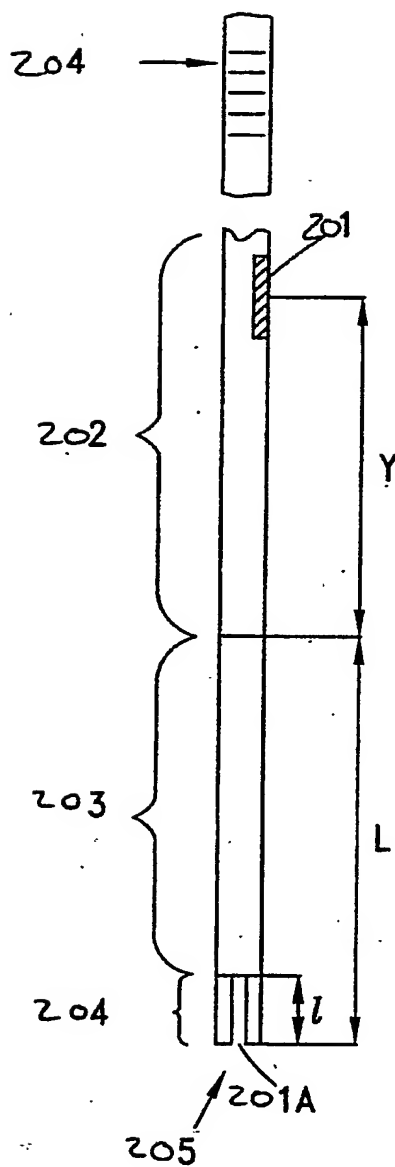


Fig. 8A

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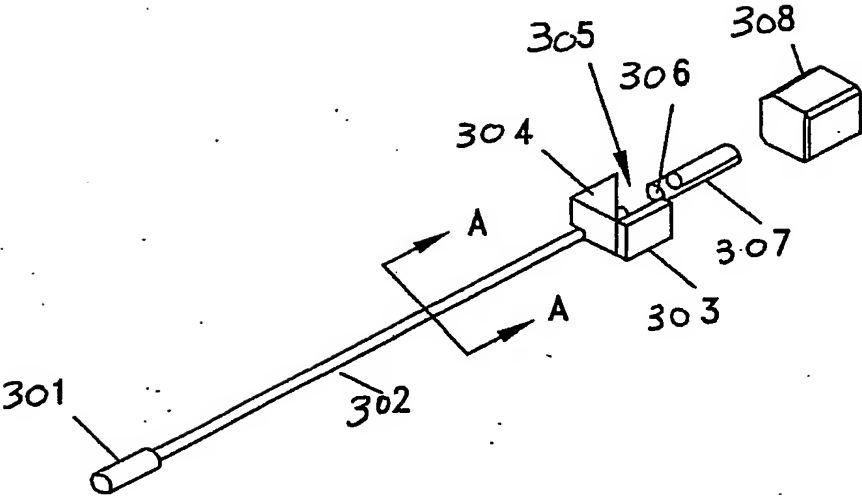


Fig. 9A

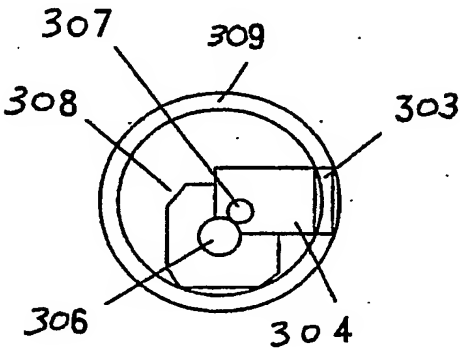


Fig. 9B

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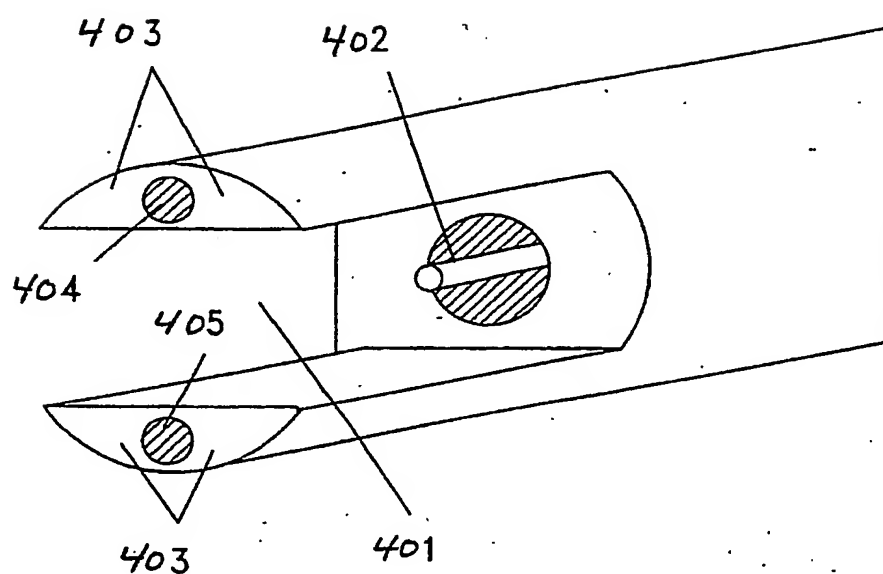


Fig. 10

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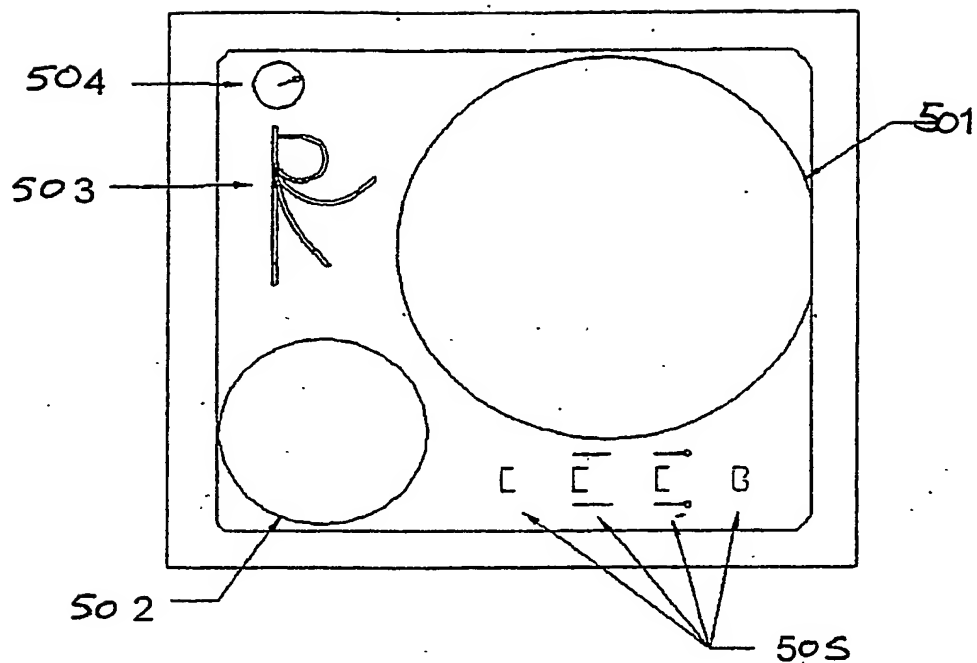


Fig. 11

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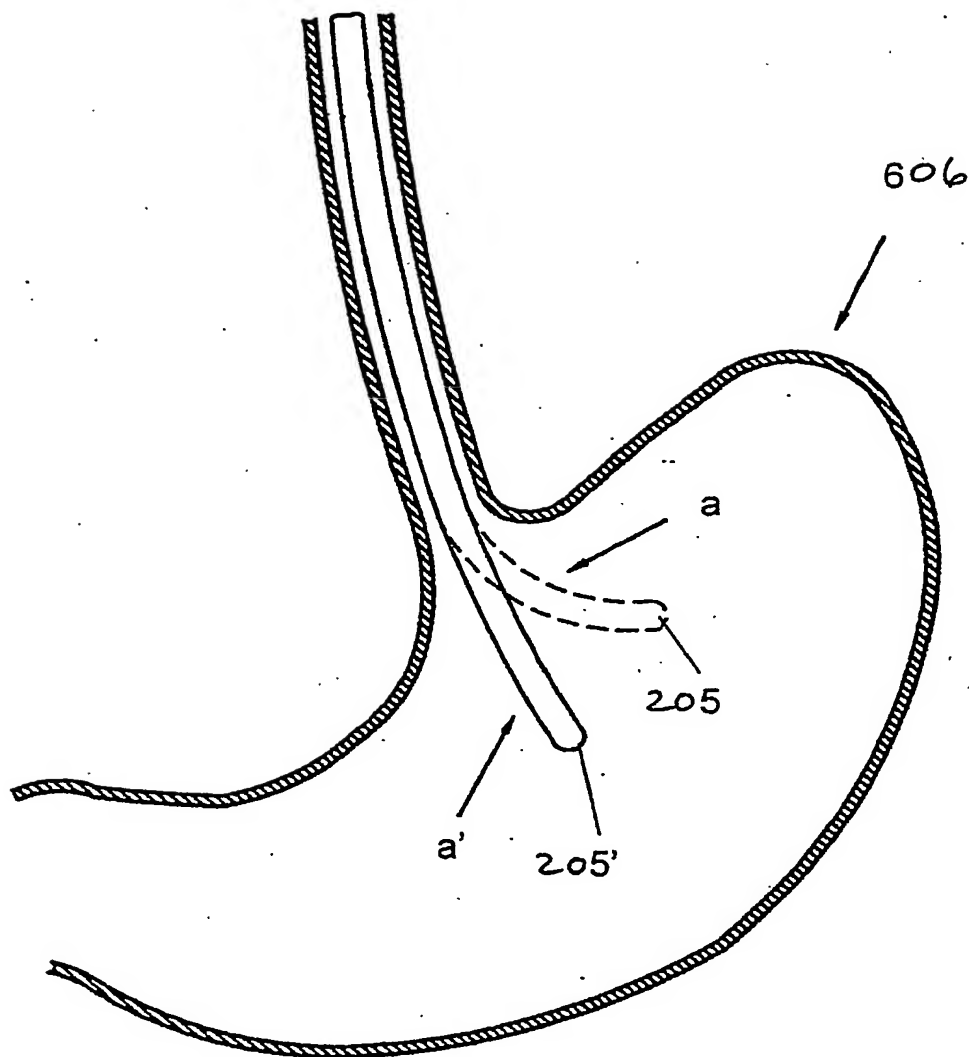


Fig. 12A



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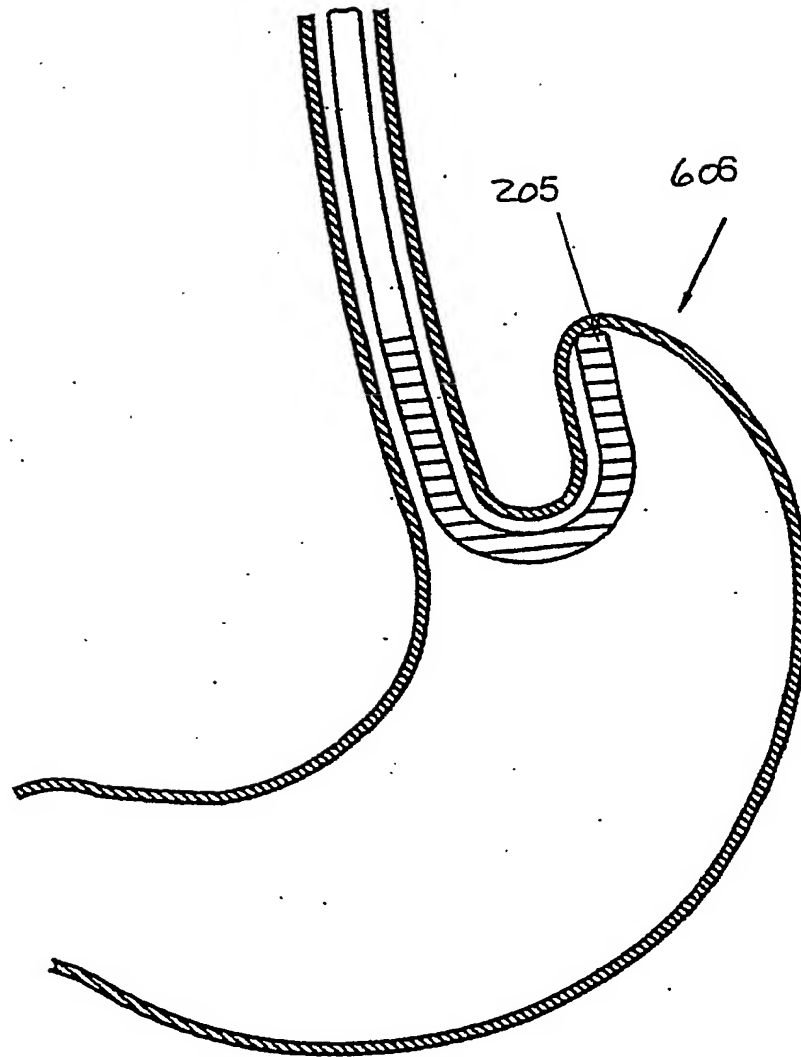


Fig. 12B

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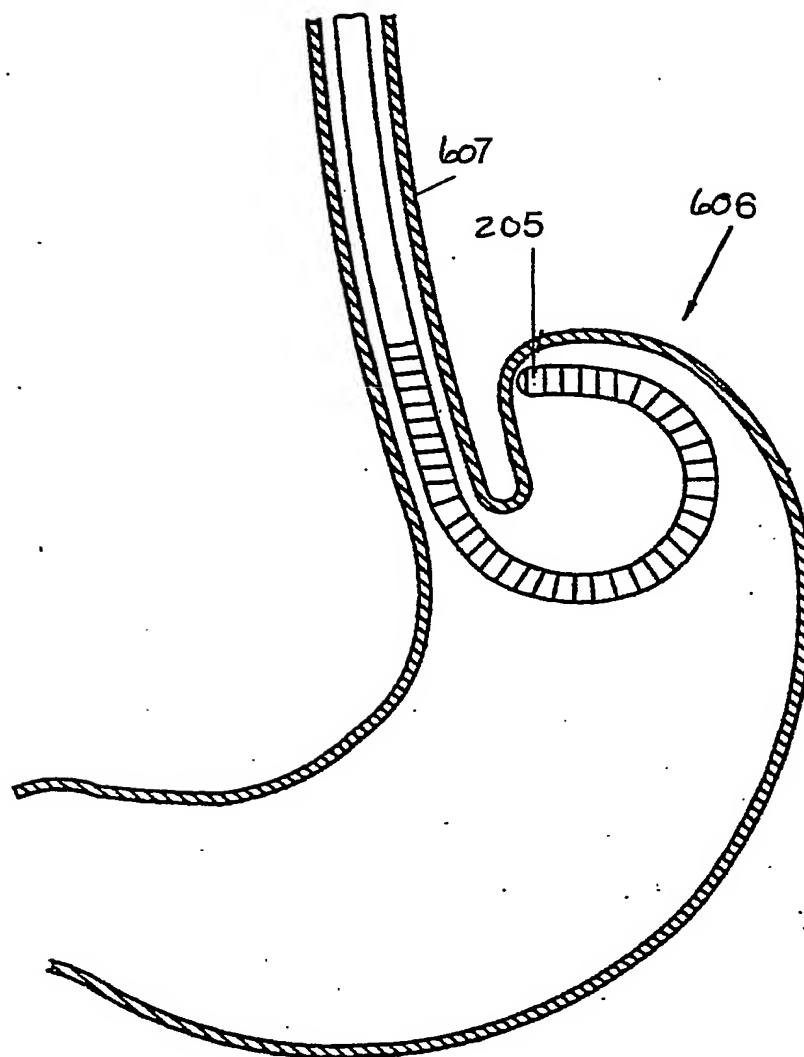


Fig. 12C

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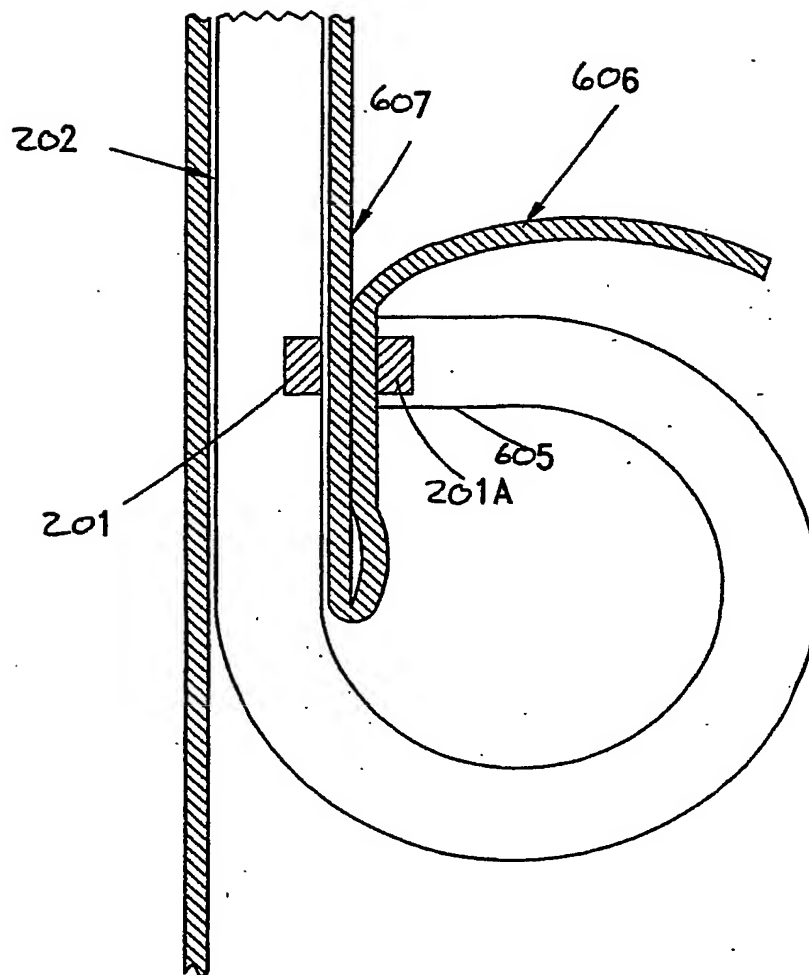


Fig. 13

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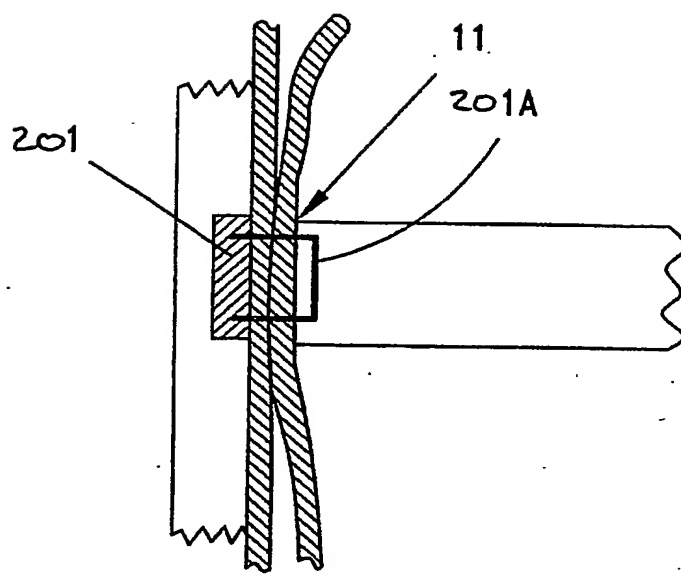


Fig. 14A

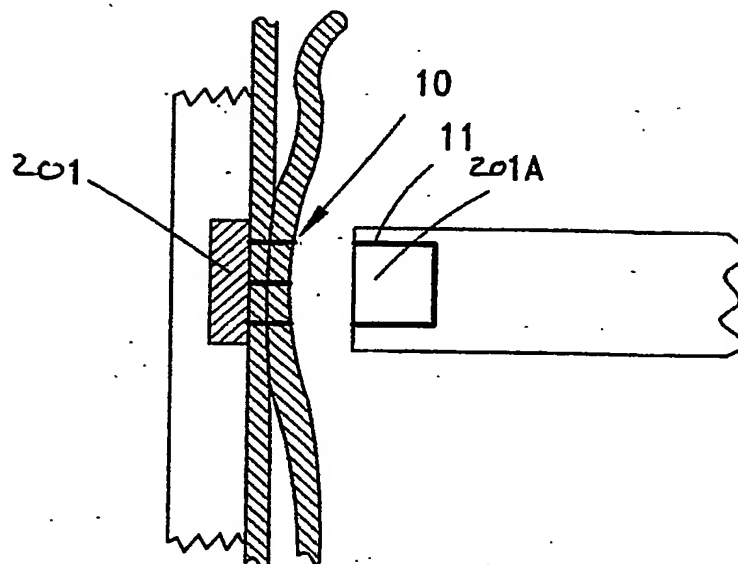


Fig. 14B